

(19) World Intellectual Property Organization
International Bureau



(43) International Publication Date
30 January 2003 (30.01.2003)

PCT

(10) International Publication Number
WO 03/008618 A2

(51) International Patent Classification⁷: **C12P 19/18**,
19/08, C12N 15/52, 9/10, 1/20, 1/21, C08B 37/02, A23L
1/054 // (C12P 19/18, C12R 1:225)

(74) Agent: **JORRITSMA, Ruurd**; Nederlandsch Octrooi-
bureau, Scheveningseweg 82, P.O. Box 29720, NL-2502 LS
The Hague (NL).

(21) International Application Number: **PCT/NL02/00495**

(81) Designated States (*national*): AE, AG, AL, AM, AT, AU,
AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU,
CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,
GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC,
LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW,
MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG,
SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ,
VN, YU, ZA, ZM, ZW.

(22) International Filing Date: **22 July 2002 (22.07.2002)**

(25) Filing Language: **English**

(26) Publication Language: **English**

(30) Priority Data:
01202752.0 20 July 2001 (20.07.2001) EP
01202841.1 25 July 2001 (25.07.2001) EP

(84) Designated States (*regional*): ARIPO patent (GH, GM,
KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW),
Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM),
European patent (AT, BE, BG, CH, CY, CZ, DE, DK, EE,
ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK,
TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GQ,
GW, ML, MR, NE, SN, TD, TG).

(71) Applicant (*for all designated States except US*): **NED-
ERLANDSE ORGANISATIE VOOR TOEGEPAST-
NATUUR-WETENSCHAPPELIJK ONDERZOEK
TNO [NL/NL]; Schoemakerstraat 97, NL-2628 VK Delft
(NL).**

Published:

— *without international search report and to be republished
upon receipt of that report*

(72) Inventor; and

(75) Inventor/Applicant (*for US only*): **VAN GEEL-SCHUT-
TEN, Gerritdina, Hendrika [NL/NL]; Bosstraat 78,
NL-3971 XG Driebergen (NL).**

*For two-letter codes and other abbreviations, refer to the "Guid-
ance Notes on Codes and Abbreviations" appearing at the begin-
ning of each regular issue of the PCT Gazette.*

(54) Title: **NOVEL GLUCANS AND NOVEL GLUCANSUCRASES DERIVED FROM LACTIC ACID BACTERIA**

(57) Abstract: The invention pertains to novel glucans capable of being produced by glucosyltransferase activity of a lactic acid bacterium on a sucrose substrate, the glucan having an average molecular weight between 10 kDa and 1 GDa, consisting essentially of $\alpha(1,3)$ - and $\alpha(1,6)$ -linked anhydroglucose units (AGU) and to glucansucrases capable of producing these glucans from sucrose. The glucans have thickening and anti-corrosive properties. The glucans can be chemically modified



WO 03/008618 A2

Novel glucans and novel glucansucrases derived from lactic acid bacteria

[0001] The present invention is in the field of enzymatic production of biomolecules. The invention is particularly concerned with novel glucans derived from lactic acid bacteria, with novel glucosyl-transferases derived from such bacteria and with a process for production of new and useful glucans from sucrose.

Background of the invention

[0002] Several bacteria are known to produce exopolysaccharides, i.e. polysaccharides secreted into the culture medium. Well-known examples of bacterial exopolysaccharides include xanthan from *Xanthomonas campestris*, gellan from *Sphingomonas paucimobilis* and pullulan from *Aureobasidium pullulans*. Lactic acid bacteria known to produce exopolysaccharides include *Leuconostoc mesenteroides* strains producing dextrans, $\alpha(1\rightarrow 6)$ -linked poly-anhydroglucose, and alternans i.e. poly-anhydroglucoses having alternating $\alpha(1\rightarrow 6)$ and $\alpha(1\rightarrow 3)$ -linkages, oral *Streptococcus* strains producing glucans responsible for dental plaque formation, and a particular *Lactobacillus reuteri* strain producing $\alpha(1,6)$ - and $\alpha(1,4)$ -linked anhydroglucose (Van Geel-Schutten, *et al.*, *Appl. Environ. Microbiol.* (1999) 65, 3008-3014). The properties of exopolysaccharides depend on the type of monosaccharide units, the type of linkages, the degree and type of branching, the length of the polysaccharide chain, the molecular weight and the conformation of the polymers.

[0003] Argüello-Morales *et al.* (*FEMS Microbiol. Lett.* 182 (2000) 81-85) describe an alternansucrase from *Leuconostoc mesenteroides* NRRL B-1355. Monchois *et al.* (*Gene* 182 (1996) 23-32; *FEMS Microbiol. Lett.* 159 (1998) 307-315) for instance describe two different dextransucrases from *Lc. mesenteroides* NRRL B-1299. A method for selecting *Leuconostoc mesenteroides* strains that produce a high proportion of alternan to dextran is described in US 5,789,209. The prior art does not disclose or suggest other lactic acid bacteria than *Leuconostoc* or *Streptococcus* that are capable of producing glucans having both $\alpha(1\rightarrow 6)$ and $\alpha(1\rightarrow 3)$ -linkages.

Summary of the invention

[0004] Several lactic acid bacteria strains were found, according to the invention, to be capable of producing a particular class of glucans. These glucans have in common that their anhydroglucose units (AGU) are linked $\alpha(1,3)$ - and/or $\alpha(1,6)$ -glucosidic bonds, i.e. they are α -glucans largely or completely devoid of $\alpha(1,4)$ -bonds. These glucans may be of

the alternan (alternating $\alpha(1,3)$ and $\alpha(1,6)$ linkages), mutan (mixed $\alpha(1,3)$ and $\alpha(1,6)$ linkages, usually $\alpha(1,3)$ predominant) or dextran (mainly $\alpha(1,6)$ linkages, some $\alpha(1,3)$) type, or other type. The glucans can be produced from sucrose, using sucrase enzymes which are active in the lactic acid bacteria. They can be produced on a large scale and
5 isolated in a commercially feasible way, as the glucans are produced outside the bacterial cell, or even in the absence of the bacteria, using isolated sucrase enzymes. The glucans are produced by food-grade strains and have interesting properties, such as prebiotic utility or thickening of water-based compositions.

[0005] The invention is concerned with these novel glucans, with the lactic acid bacterial,
10 especially *Lactobacillus* strains and their enzymic proteins that produce these glucans from sucrose, as well as with methods for producing the glucans using the strains and/or their enzymes, with nucleotide sequences encoding these enzymic proteins which convert sucrose, with the use of the glucans as thickeners, prebiotics, anticorrosives, etc., and as starting materials for modified glucans.

15 **Description of the invention**

[0006] The invention pertains to *Lactobacillus* strains containing a glucosyltransferase (glucansucrase) capable of producing a glucan having at least 10 anhydroglucose units (AGU) having a backbone consisting essentially of $\alpha(1,3)$ - and/or $\alpha(1,6)$ -linked AGU, in the presence of sucrose. Such strains can be found among current sources of *Lactobacilli*,
20 such as food sources, silage, mammalian samples etc. These strains containing the glucosyltransferases and producing the glucans can be identified by isolating *Lactobacillus* strains from these sources, growing them on sucrose and analysing the polysaccharide product using suitable analytical methods such as chromatography. The genes encoding these glucosyltransferases can be identified by amplifying nucleotide
25 sequence fragments of the strain using primers based on known glucosyltransferase genes and retaining the positive strains (see examples). Several glucan-producing strains were isolated and identified from different sources and different *Lactobacillus* species, such as *Lb. reuteri*, *Lb. fermentum*, *Lb. sake* and *Lb. parabuechneri* or related species. The glucosyltransferases from these glucan-producing strains were also identified and,
30 completely or partly, sequenced (see Examples).

[0007] The novel glucans of the invention are capable of being produced by glucosyltransferase (glucansucrase) activity of a lactic acid bacterium on a sucrose donor substrate. The glucans have an average molecular weight between 10 kDa and 1 GDa, and

consist essentially of $\alpha(1,3)$ - and/or $\alpha(1,6)$ -linked anhydroglucose units (AGU), to which side-chains also consisting of $\alpha(1,3)$ - and/or $\alpha(1,6)$ -linked AGU may be attached.

[0008] In particular, the glucans according to the invention either comprise 15-80% of $\alpha(1,3)$ -linked AGU, 2-80%, especially 4-80% and more especially 15-80% of $\alpha(1,6)$ -linked and 2-25% of $\alpha(1,3,6)$ -linked (branching) AGU, or 80-99% of $\alpha(1,6)$ -linked AGU and 1-20% of $\alpha(1,3)$ -linked or $\alpha(1,3,6)$ -linked (branching) AGU, in particular 1-15% of $\alpha(1,3)$ -linked AGU and 5-15% of $\alpha(1,3)$ - and $\alpha(1,3,6)$ -linked units taken together. Thus, the invention covers a glucan having an average molecular weight of 50 kDa to 1 MDa and comprising 25-50%, especially 29-39% of $\alpha(1,3)$ -linked AGU, 20-45%, especially 30-40% of $\alpha(1,6)$ -linked AGU, 5-25%, especially 3-13% of $\alpha(1,3,6)$ -linked AGU and 6-30% of terminal AGU. Furthermore, the invention pertains to a glucan having an average molecular weight of 10-50 MDa and comprising 15-26% $\alpha(1,3)$ -linked AGU, 30-50% of $\alpha(1,6)$ -linked AGU, 5-20% of $\alpha(1,3,6)$ -linked AGU and 5-35% of terminal AGU. Also, in another embodiment the invention covers a glucan having an average molecular weight of 1-50 MDa and comprising 40-60% of $\alpha(1,3)$ -linked AGU, 2-20%, especially 2-12% of $\alpha(1,6)$ -linked AGU, 10-25% of $\alpha(1,3,6)$ -linked AGU and 10-30% of terminal AGU. In yet another embodiment, the invention comprises a glucan having an average molecular weight of 10-50 MDa and comprising 80-99%, especially 88-99% and more especially 90-99% of $\alpha(1,6)$ -linked AGU, or 80-90% of $\alpha(1,6)$ - and 1-10% of $\alpha(1,3)$ -linked AGU, the remainder being 1,3,6 linked and terminal AGU.

[0009] The invention also concerns the enzymes originating from lactic acid bacteria, or from recombinant sources, capable of producing the glucans described above starting from sucrose. The enzymes are new and they can be classified as glucansucrases or glucosyltransferases. Their partial sequence information is given below in SEQ ID No's 1-10. More complete sequence information is given in SEQ ID No's 11-22. Proteins according to the invention comprise an amino acid sequence exhibiting at least 70%, preferably at least 80%, most preferably at least 90%, amino acid identity with any one of the amino acid sequences of SEQ ID No. 2, 4, 8, 10, 12, 14, 16, 18, 20 and 22 or of stretches of at least 221-224 amino acids thereof, or at least 100 contiguous amino acids exhibiting at least 80%, preferably at least 90%, amino acid identity with these sequences. Further preferred sequences are indicated in the description of the alignment figure given below.

[0010] The enzymes can be used as such for producing the glucans described above, or for producing oligosaccharides and polysaccharides having a similar $\alpha(1,3)$ and/or $\alpha(1,6)$ linked glucan structure. Their genes can also be incorporated in suitable host organisms, to produce alternative glucan-production systems. The invention also pertains to such
5 recombinant, preferably food-grade microorganisms, e.g. bacteria, especially lactic acid bacteria, yeasts, fungi etc., containing the genes of the glucansucrases described above and being capable of expressing the glucansucrases.

[0011] The invention also pertains to a process of producing a glucan as described above. This glucan can be produced by a *Lactobacillus* strain as described above, or by a
10 recombinant micro-organism expressing the glucosyltransferase according to the invention or by an isolated glucosyltransferase according to the invention and a suitable glucose source such as for instance sucrose. The glucosyltransferase may be isolated by conventional means from the culture of a glucosyltransferase-positive lactic acid bacterium, especially a *Lactobacillus* species, or from a recombinant organism expressing
15 the glucosyltransferase gene.

[0012] The glucan and the gluco-oligosaccharides produced by the *Lactobacillus* strains can be recovered from the culture supernatant of *Lactobacillus* strains described above, containing the glucosyltransferase according to the invention. The glucan can comprise at least 20, up to about 100,000 α -anhydroglucose units with the unique structure described
20 above.

[0013] The glucan-producing enzymes according to invention, or at least the most preferred ones, are constitutive in the *Lactobacillus* strains, in that they are always present. This is contrast to most glucan (dextran-) producing *Leuconostoc* strains of the prior art, wherein the enzymes are only expressed upon growth in the presence of sucrose.
25 This allows a more efficient production of glucans by the microorganisms of the invention.

[0014] The glucans according to invention have a variety of useful properties. They are suitable as prebiotics, and thus they can be incorporated in nutritional or pharmaceutical compositions intended for improving the condition of the gastrointestinal tract. For this
30 purpose, they can be used as such or in the form of their oligosaccharides. They can also be combined with other poly- or oligosaccharides, such as fructans, galactans, xylans, arabinans, mannans, indigestible glucans and hetero-oligosaccharides, or with probiotic micro-organisms, including the lactic acid bacteria from which the glucans originate, resulting in synbiotic compositions. The glucans and their shortened homologues are also

useful as bioactive agents, e.g. as immunomodulators, anti-ulcer agents and cholesterol-lowering agents.

[0015] The glucans are also useful as thickening agents. As such they can be incorporated in foodstuffs such as beverages, sauces, dressings, dairy products, in amounts of from 1 g/l to about 100 g/l, especially about 10 to 50 g/l.

[0016] The glucans of the invention are furthermore useful as anticorrosion agents, for example for the protection of ship hulls. For that purpose, they may be applied in the form of solutions or suspensions, by spraying, coating, dipping and other techniques known in the art of corrosion control.

[0017] The glucans can be used as such. They can also be modified by physical or chemical means. Suitable examples of chemical modification include oxidation, especially 2,3- or 3,4-oxidation using periodate or hypohalite, in glucans having α -1,6 linkages, or 6-oxidation using nitroxyls with peracid or hypohalite in glucans having α -1,3 linkages. Hypohalite oxidation resulting in ring-opened 2,3- or 3,4-dicarboxy-anhydroglucose units (see e.g. EP-A-427349), while periodate oxidation results in ring-opened 2,3- or 3,4-dialdehyde-anhydroglucose units (see e.g. WO 95/12619), which can be further oxidised to (partially) carboxylated units (see e.g. WO 00/26257). Nitroxyl-mediated oxidation using hypochlorite or a peracid results in 6-aldehyde- and 6-carboxy-anhydroglucose units (see e.g. WO 95/07303).

[0018] The oxidised glucans have improved water-solubility, altered viscosity and a retarded fermentability and can be used as metal-complexing agents, detergent additives, strengthening additives, bioactive carbohydrates, emulsifiers and water binding agents. They can also be used as starting materials for further derivatisation such as cross-linking and the introduction of hydrophobes. Oxidised glucans coupled to proteins can be used as emulsifiers and stabilisers. The oxidised glucans of the invention preferably contain 0.05-1.0 carboxyl groups, more preferably 0.2-0.8 carboxyl groups per anhydroglucose unit, e.g. as 6-carboxyl groups on 1,3-linked units.

[0019] When modified glucans with high proportion of carboxyl groups are desired, two oxidation processes can be combined or an oxidation can be combined with e.g. carboxymethylation (see below). Thus, an α -(1,3/1,6)-glucan having a degree of substitution (DS) for carboxyl groups between 0,3 and 1,0 can be conveniently prepared by first nitroxyl-mediated oxidation, resulting in 1,3-substituted units being oxidation to glucuronic acid units, followed by e.g. periodate and chlorite oxidation, resulting in 1,6-substituted units* being converted to ring-opened dicarboxy-substituted units. The order

of processes can also be inverted, or one oxidation process, such as nitroxyl-mediated 6-oxidation can be combined with carboxymethylation. Also, by appropriate adaptation of the oxidation processes mixed aldehyde-containing and carboxyl-containing polymers can be obtained.

- 5 [0020] Other useful modifications are alkylation, acylation, hydroxyalkylation, amino-alkylation, carboxyalkylation, phosphorylation, sulphatation, as well as physical and chemical crosslinking. Phosphorylation (see: O.B. Wurzburg (1986), Modified Starches: properties and uses. CRC Press Inc., Boca Raton, 97-112) can be achieved by dry heating glucans with a mixture of monosodium and disodium hydrogen phosphate or with tripoly-
10 phosphate. The phosphorylated glucans are suitable as wet-end additives in papermaking, as binders in paper coating compositions, as warp sizing-agents, and as core binders for sand molds for metal casting. Acylation, especially acetylation or propionylation using acetic or propionic anhydride respectively, results in products suitable as bleaching assistants and for the use in foils. Acylation with e.g. alkenyl succinic anhydrides or
15 (activated) fatty acids results in surface-active products suitable as e.g. surfactants, emulsifiers, and stabilisers. Crosslinking, e.g. by coupling oxidised derivatives, or by reaction with a crosslinking agent such as triphosphoric acid, epichlorohydrine or a dialdehyde, can be used to adjust the physical properties of the glucans, e.g. to enhance their water-binding or thickening capacities.
- 20 [0021] Hydroxyalkylation is commonly performed by base-catalysed reaction with alkylene oxides, such as ethylene oxide, propylene oxide or epichlorohydrin; the hydroxy-alkylated products have improved solubility and viscosity characteristics. Carboxy-methylation is achieved by reaction of the glucans with monochloroacetic acid or its alkali metal salts and results in anionic polymers suitable for various purposes including
25 crystallisation inhibitors, and metal complexants. Amino-alkylation can be achieved by reaction of the glucans with alkylene-imines, halo-alkyl amines or amino-alkylene oxides, or by reaction of epichlorohydrine adducts of the glucans with suitable amines. These products can be used as cationic polymers in a variety of applications, especially as a wet-end additive in paper making to increase strength, for filler and fines retention, and to
30 improve the drainage rate of paper pulp. Other potential applications include textile sizing and wastewater purification. The above mentioned modifications can be used either separately or in combination depending on the desired product. Furthermore, the degree of chemical modification is variable and depends on the intended use. If necessary 100% modification, i.e. modification of all anhydroglucose units can be performed. However,

partial modification, e.g. from less than 1 (e.g. 0.2) modified anhydroglucose unit per 100 units up to higher levels, will often be sufficient in order to obtain the desired effect.

[0022] Another suitable type of derivatives is formed by hydrolysates of the present glucans. Hydrolysis can be performed in a controlled manner in a way known per se, using e.g. dilute acid or glucanolytic enzymes, especially α -1,3-glucanases or α -1,6 glucanases. Hydrolysis results in polysaccharides of reduced chain length (degree of polymerisation, DP, of more than 20) or oligosaccharides (DP of less than 20).

[0023] The invention also relates to gluco-oligosaccharides containing the characteristic structure of the glucan described above. These can be produced using an isolated glucan-sucrase according to the invention or a *Lactobacillus* strain, or a recombinant micro-organism containing (a part of) a glucosyltransferase according to the invention. Gluco-oligosaccharides thus produced can be used as prebiotics and probiotics. The production of the gluco-oligosaccharides is different from the glucan synthesis reaction. In addition to sucrose, the substrate of the glucansucrase, an acceptor molecule such as maltose or lactose can be used as an acceptor, to synthesise oligosaccharides. Consecutive attachment of glucose units in a manner determined by the particular glucansucrase results in α (1,3)- and/or α (1,6)-linked gluco-oligosaccharides, the chain length of which can be determined by selecting the appropriate reaction conditions. Longer reaction times, higher sucrose levels and lower acceptor levels will usually result in relatively long chains, e.g. having a degree of polymerisation (DP) of more than 10, up to several hundreds if desired, while shorter reaction times, lower sucrose levels and higher acceptor levels will result in relatively short chains, e.g. with a DP from about 3 up to 10 or higher. Another way of producing gluco-oligosaccharides is by hydrolysis of the glucan described above. This hydrolysis can be performed by known hydrolysis methods such as enzymatic hydrolysis with enzymes such as amylase, dextranase or pullulanase or by acid hydrolysis. The produced gluco-oligosaccharides contain at least one 1,6- or one 1,3-glucosidic link to be used as prebiotics.

[0024] The invention also relates to a probiotic or synbiotic composition containing a *Lactobacillus* strain capable of producing a glucan and/or gluco-oligosaccharide according to the invention. The strain may also produce another poorly digestible poly- or oligosaccharide, such as a fructan. The probiotic or synbiotic compositions of the invention may be directly ingested with or without a suitable vehicle or used as an additive in conjunction with foods. They can be incorporated into a variety of foods and beverages including, but not limited to, yoghurts, ice creams, cheeses, baked products

such as bread, biscuits and cakes, dairy and dairy substitute foods, confectionery products, edible oil compositions, spreads, breakfast cereals, juices and the like.

[0025] Furthermore, the invention pertains to a process of improving the microbial status in the mammalian colon comprising administering an effective amount of a *Lactobacillus* strain capable of producing a glucan and/or gluco-oligosaccharide according to the invention. Furthermore, a process of improving the microbial status of the mammalian colon comprising administering an effective amount of a glucan or gluco-oligosaccharide according to the invention is also a part of the present invention.

10 **Examples**

General

The various lactic acid bacterial strains were isolated from a variety of sources, including fermented foods, the gastrointestinal tract of various human or animal species, and silage.

15 **Example 1: Identification and nucleotide sequence of glucansucrase/glucosyltransferase genes from lactobacilli**

The glucansucrase genes were identified by amplification with PCR using degenerated primers (GTFrev, 5' ADRTC NCCRT ARTAN AVNYK NG 3' and GTFforw, 5'-GAYAA YWSNA AYCCNRYNGT NC-3'; N = A, C, G or T, Y = T or C, K = G or T, W = A or T, S = C or G, R = A or G), based on conserved amino acid sequences of different published glucansucrase genes. An amplification product with the predicted size of about 660 bp was obtained and cloned in *Escherichia coli* Top 10 using pCR-XL-TOPO (Invitrogen). Sequence analysis confirmed that part of a *gtf* gene had been isolated. The 660 bp amplified was used to design primers for inversed PCR. For inverse PCR chromosomal DNA was digested with 10 different enzymes ligated, yielding circular DNA molecules. PCR with the diverging primers with the circular ligation products as template yielded amplicons of various sizes, those products were cloned into pCR-XL-TOPO (Invitrogen) and sequenced (GATC, Konstanz, Germany). If necessary additional inverse PCR reactions were carried out to obtain the complete gene(s). Both strands of the entire glucansucrase genes were sequenced twice.

30 **Example 2: Isolation and identification of α -(1,6) glucan and a glucansucrase from *Lactobacillus reuteri* strain 180**

L. reuteri strain 180 was deposited as LMG P-18389 at the BCCM/LMG Culture Collection at Gent, Belgium. The strain was grown in 18 litres of MRS-s medium (in g per kg): yeast extract (22), sodium acetate trihydrate (5), sodium citrate dihydrate (2.42), ammonium chloride (1.32), dipotassium hydrogen phosphate (2), magnesium sulphate heptahydrate (0.2), manganese sulphate heptahydrate (0.05), sorbitan mono-oleate (1), vitamins (in mg per kg: B1: 14.4, B2: 3.6, B3: 72, H 0.216), sucrose (100), tap water

(remainder), for 21 h at 37°C under anaerobic conditions (pH 5.5). See also: Van Geel-Schutten et al., Appl. Microbiol. Biotechnol. (1998) 50, 697-703. During growth, 13 g/l polysaccharide was produced. This polysaccharide was isolated as described in the reference cited above. The monosaccharide composition of the polysaccharide was determined by hydrolysis of the soluble part of the polysaccharide and high-performance anion-exchange chromatography. It was characterised as a glucan. This glucan was not formed when the strain was grown on glucose instead of sucrose. Methylation analysis (Van Geel-Schutten et al. 1999) revealed the presence of 17-24% $\alpha(1,3)$ -linked glucosyl units, 34-44% of $\alpha(1,6)$ -linked glucosyl units, 7-15% of $\alpha(1,3,6)$ -linked glucosyl units and 7-35% of terminal glucosyl units. The average molecular weight of the glucan was determined to be 3.6×10^7 Da and the Rg was 45 nm.

The average molecular weight of the polysaccharide was established using the SEC-MALLS system: 0.0522 g of the glucan was dissolved in 10 ml DMSO/water (90/10) and heated for 1 hour at 80°C, filtered through a 0.45 μ m filter and injected on the SEC-MALLS system and analysed using the following conditions:

Eluent:	DMSO/water (90/10) with 0.1 M NaNO ₃
Flow rate:	0.5 ml/min
Injection volume:	0.247 ml
Column:	PLgel Guard, mixed-A and mixed-D
Temperature:	90°C

Detection: MALLS (DAWN-DSP), 50°C, A₂=0, dn/dc=0.074, F2 cell, RI; SDS PAGE followed by PAS-staining (Van Geel-Schutten et al. 1999) revealed the presence of an extracellular sucrase with a molecular weight of about 190 kDa. Part of the gene encoding the sucrase enzyme was isolated using PCR techniques and sequenced. On the deduced amino acid sequence of the fragment, high homologies were found with other glucan-sucrases. This partial sequence information is given in SEQ ID No. 1 (DNA) and 2 (protein). Full sequence information is given in SEQ ID No's. 11 and 12.

The glucan produced by *L. reuteri* strain 180 has been tested for application on ship hulls for the prevention of corrosion (see Example 8).

30

Example 3: Isolation and identification of $\alpha(1,6/1,3)$ glucan and a glucansucrase from Lactobacillus reuteri strain ML1

L. reuteri strain ML1, deposited as LMG P-20347 at the BCCM/LMG Culture Collection at Gent, Belgium, was grown overnight under anaerobic conditions at 37°C on MRS supplemented with sucrose (see Example 2). The cells were removed by centrifugation and two volumes of ethanol were added to the supernatant. The precipitated polysaccharides were harvested by centrifugation and resuspended in 2-3 liters of demi water and precipitated again with two volumes of ethanol. The glucan produced by this strain (7 g) was characterised by methylation analysis and monosaccharide composition analysis as

described in Example 2. The polymer was found to consist of 48-53% of $\alpha(1-3)$ linked glucosyl units, 3-8% of $\alpha(1-6)$ linked glucosyl units, 12-20% of $\alpha(1-3-6)$ linked glucosyl units (branching units) and 20-30% of 1-linked (terminal) glucose units. The glucans were not produced during growth on glucose. The average molecular weight of the polysaccharide was established to be 7.6×10^6 Da using the SEC-MALLS system as described in example 2. These were the first examples of the production of mutan-like polymers by lactobacilli. The glucan produced by *L. reuteri* strain ML1 has been tested for application as anticorrosive agent and showed excellent utility for the prevention of corrosion e.g. on ship hulls.

SDS PAGE followed by PAS-staining (Van Geel-Schutten et al. 1999) revealed the presence of an extracellular sucrase with a molecular weight of about 190 kDa. It was found that this strain produces two glucansucrases. Sequence information for these sucrase is given in SEQ ID No's 13 and 14 (ML1) and 15 and 16 (ML4).

Example 4: Isolation and identification of $\alpha(1,6/1,3)$ glucan and a glucansucrase from *Lactobacillus* strain LB 33.

A new *Lactobacillus* strain was obtained and was deposited as LMG P-20349. The strain was identified by 16S rRNA to be most closely related to *Lactobacillus parabuchneri*. The strain grown overnight on MRS supplemented with sucrose under anaerobic conditions at 37°C (see Example 2). 420 gram of glucan was produced. The glucan produced by this strain is not produced during growth on glucose.

Methylation analysis (see Example 2) revealed that the polymer consists of equal amounts of 29-39% of $\alpha(1-3)$ linked glucosyl units, 30-40% of $\alpha(1-6)$ linked glucosyl units, 3-13% of $\alpha(1-3-6)$ linked glucosyl units (branching units) and 15-30% of 1-linked (terminal) glucose units.

The average molecular weight of the polysaccharide was established to be 2×10^5 Da, using the SEC-MALLS system as described in Example 2.

By PCR with degenerated primers part of a sucrase type of glucosyl-transferase could be isolated indicating that the glucan is produced by a sucrase. This confirms the result that the glucan is produced during growth on sucrose and not on glucose. Part of the sucrase encoding gene was sequenced. On the deduced amino acid level high homologies were found with alternan sucrase from *Leuconostoc mesenteroides*. This indicates that the enzyme responsible for the glucan synthesis in *L. brevis* is the first alternan sucrase found in other bacteria than *Leuconostoc*. This partial sequence information is given in SEQ ID No. 3 (DNA) and 4 (protein). Full sequence information is given in SEQ ID No's. 17 and 18, respectively.

The glucan produced by this strain has thickening properties.

Example 5: Isolation and identification of α -(1,6) glucan and a glucansucrase from *Leuconostoc* strain 86

A new strain was obtained from silage and was deposited as LMG P-20350. The strain was identified by 16S rRNA to be a new *Leuconostoc* strain, most closely related to *Leuconostoc citreum*. The strain grown overnight on MRS supplemented with sucrose under anaerobic conditions at 37°C (see Example 2). 416 gram of glucan was produced. Methylation analysis of the glucan obtained revealed that more than 90 % of the glucose units was linked through an α (1,6) bond, identifying the polysaccharide as a dextran. The molecular weight of the glucan (determined as described in Example 2) was $3-4 \times 10^7$ Da and the Rg was 40 nm. The glucan is not produced during growth on glucose.

By PCR with degenerated primers 3 different fragments with part of a sucrase type of glucosyl-transferase could be isolated indicating that the glucan is produced by a sucrase and that possibly 3 sucraes are present in this strain. This confirms the result that the glucan is produced during growth on sucrose and not on glucose. Part of the sucrase encoding gene was sequenced. On the deduced amino acid level high homologies were found with DSRC and DSRB (fragment 1), alternan sucrase (fragment 2) and DSRA (fragment 3) from *Leuconostoc mesenteroides*. The sequence information is given in SEQ ID No's 5-10. *Leuconostoc citreum*, to which this new strain is most closely related, is not reported to produce dextran. The glucan produced by strain 86 has thickening properties.

Example 6: Identification of α -(1,6/1,3) glucan and a glucansucrase from *Lactobacillus sake* KG 15

Strain KG 15 was obtained from silage and was deposited as LMG P-21583. It was identified by 16S rRNA as *L. sake*. The strain was grown and the polysaccharide was recovered as described in example 2. The molecular weight of the polysaccharide was determined to be $4,7 \times 10^7$ Da (SEC MALLS) and the Rg was 92 nm. Methylation analysis (GC) revealed that the glucan produced by this strain is a largely linear dextran containing 4 % terminal glucose units, 86% of α (1,6) linked glucosyl units, 2% of α (1,3) linked glucosyl units and 8% α (1,3,6) disubstituted glucose units (branching points). The glucansucrase of this strain was sequenced (see SEQ ID No. 19 and 20).

Example 7: Identification of α -(1,6/1,3) glucan and a glucansucrase from *Lactobacillus fermentum* KG 3

Strain KG 3 was obtained from silage and was deposited as LMG P-21584. It was identified by 16S rRNA as *L. fermentum*. The strain was grown and the polysaccharide was recovered as described in example 2. The molecular weight of the polysaccharide was determined to be $2,4 \times 10^7$ Da (SEC MALLS) and the Rg was 107-119 nm. Methylation analysis (GC) revealed that the glucan produced by this strain is a largely linear dextran containing 3% terminal glucose units, 84% of α (1,6) linked glucosyl units,

8% of α (1,3) linked glucosyl units and 5% α (1,3,6) disubstituted glucose units (branching points). The glucansucrase of this strain was sequenced (SEQ ID No's 21 and 22).

5 **Example 8: Anticorrosion properties of glucans**

Plain carbon steel sheets of 1 cm² embedded in an epoxy matrix were exposed to a slightly corrosive medium (150 ml of 0.1 M LiClO₄) with or without the addition of a bacterial polysaccharide (0.2 g) for several days. The sheets were then examined visually and electrochemically from time to time. The corrosion potential (E_{corr} in mV with
10 reference to Ag/AgCl) and polarisation resistance (R_p in k Ω /cm²) are both a measure of the anti-corrosion effect. After an initial adaptation of 3-10 hours, these parameters attained a stable value. The experiments were carried with a heteropolysaccharide from *Lactobacillus sake*, and a homopolysaccharide of the invention (from LB 180 according to example 4), as well as without polysaccharide. The results are summarised in the table
15 below. It follows that the anti-corrosion properties of the glucan of the invention are superior. It was found that the homopolysaccharide of ML 1 (example 3) has at least equal anticorrosion performance as the LB 180 polysaccharide.

Table: Corrosion experiments

organism	type of poly-saccharide	aspect of treated sheet	E_{corr} (mV vs. Ag/AgCl)	R_p (k Ω /cm ²)
control	-	corrosion	-700	1.5
<i>Lb. sake</i>	hetero-polysaccharide	localised corrosion	-600	4.5
<i>Lb. 180</i>	α -glucan	thin black layer	-200	70

Example 9: Modification of α -1,3/1,6-glucan by oxidation

20 One gram (6.15 mmol of anhydroglucose units) of the α -1,3/1,6-glucan produced by strain LB 33 (example 4) is resuspended in 100 ml water. Next, 2,2,6,6-tetramethyl-piperidine-1-oxyl (TEMPO; 0.01 g, 0.065 mmol) and sodium bromide (100 mg, 1 mmol) are added and the suspension is cooled to 0°C. The reaction can also be performed without bromide. A solution of hypochlorite (3 ml, 15% solution, 6.3 mmol) of pH 10.0
25 (0°C) is added. The pH is kept constant by addition of 0.1M NaOH. After 1 hr, the solution is poured into 150 ml 96% ethanol, causing the product to precipitate. The white precipitate is centrifuged, resuspended in ethanol/water (70/30 v/v) and centrifuged again. Next, the precipitate is resuspended in 96% ethanol, centrifuged and dried. The uronic acid content is determined by means of the uronic acid assay according to Blumenkrantz
30 and Abdoe-Hansen (*Anal. Biochem.* 54 (1973), 484). A calibration curve was generated using polygalacturonic acid (5, 10, 15 and 20 μ g). With this calibration curve the uronic

acid content in a sample of 20 µg of the product is determined. The major part of 6-hydroxyl groups have been oxidised to carboxyl groups.

Example 10: Construction of plasmids for expression of the glucansucrase genes in *E. coli*.

5 Two primers were designed with appropriate restriction sites; the C-terminal primer contained in all cases a His-tag. The PCR products were first cloned in pCR-XL-TOPO. The PCR products were removed from pCR-XL-TOPO using the appropriate enzymes and ligated in the appropriate sites of an expression vector (e.g pET15b (Novagen)).

For the expression of part of the glucosyltransferase gene of LB 180 (for better
10 expression, the N-terminal region encoding the N-terminal variable domain of the glucansucrase, was not cloned) in *E. coli*, a PCR reaction was performed using Forw180 (5'-GATGCATGAG **CTCCCATGGG** CATTAACGGC CAACAATATT ATTATTGACC C-3') containing *SacI* (bold) and *NcoI* (underlined) sites, and Rev180 (5'-ATATCGATGG **GCCCCGGATC** CTATTAGTGA *TGGTGATGGT GATGTTTTTG*
15 *GCCGTTTAAA TCACCAGGTT TTAATGG*-3'), containing *ApaI* (bold), *BamHI* (underlined) and a 6x His-tag (italics) as primers. The PCR product was cloned in pCR-XL-TOPO. The PCR product was removed from pCR-XL-TOPO using *NcoI/BamHI* and ligated in the corresponding sites of pET15b (Novagen). The resulting plasmid (pET15b180) containing part of the glucansucrase gene of 704 amino acids encoding a
20 glucansucrase without the variable N-terminal domain was transformed to *E. coli* B121 DE3 star (Invitrogen).

Cells of *E. coli* harbouring the pET15b180 were harvested by centrifugation after 16 h of growth under aerobic conditions at 37 °C. The pellet was washed with 50 mM sodium acetate buffer pH 5.5 containing 1 mM CaCl₂ and 1% (v/v) Tween 80 and the suspension
25 was centrifuged again. Pelleted cells were resuspended in with 50 mM sodium acetate buffer pH 5.5 containing 1 mM CaCl₂ and 1% (v/v) Tween 80, and 7.2 mM β-mercaptoethanol. Cells were broken by sonication and cell debris and intact cells were removed by centrifugation for 15 minutes at 4 °C at 14,000 rpm (Eppendorf). The resulting cell free extract was used as enzyme source to produce high molecular weight glucans from
30 sucrose in 50 mM sodium acetate buffer pH 5.5 containing 1 mM CaCl₂ and 1% (v/v) Tween 80 and 10 g/l sucrose. After 16 hours of incubation, the glucans were isolated using ethanol precipitation. When cell free extracts of *E. coli* B121 DE3 star (Invitrogen) harbouring the plasmid pET15b (without insert) were used as enzyme source, no glucans were produced from sucrose.

35

Sequence information

SEQ ID No's 1 and 2 give the nucleotide and amino acid sequence, respectively, of a part of the glucansucrase from strain Lb180 as originally determined (Example 2). The partial

sequence shows 53% (199/223) sequence identity and 68% similarity with dextransucrase DSRB742 of *Leuconostoc mesenteroides* (*Lc. mes.*), with 2 gaps (between amino acids F172 and N173), and 52% identity with some other dextransucrases and alternansucrases of *Lc. mes.*

- 5 SEQ ID No's 3 and 4 give the nucleotide and amino acid sequence, respectively, of a part of the glucansucrase from strain Lb 33 as originally determined (Example 4). The partial sequence shows 63% (143/224) sequence identity and 75% similarity with dextransucrase DSRB742 of *Lc. mes.* with 1 gap.

- 10 SEQ ID No's 5 and 6 give the nucleotide and amino acid sequence, respectively, of a part of a glucansucrase (86-1) from strain Lc 86 (Example 5). The partial sequence shows 98% (219/223) sequence identity and 99% similarity with dextransucrase DSRB742 of *Lc. mes.*

- 15 SEQ ID No's 7 and 8 give the nucleotide and amino acid sequence, respectively, of a part of another glucansucrase (86-5) from strain Lc 86 (Example 5). The partial sequence shows 55% (123/223) sequence identity and 68% similarity with dextransucrase DSRB742 of *Lc. mes.*, with 2 gaps (between amino acids M128 and R129 and between D162 and H163), and 51-56% identity with some other dextransucrases and alternansucrases of *Lc. mes.*

- 20 SEQ ID No's 9 and 10 give the nucleotide and amino acid sequence, respectively, of another glucansucrase (86-8) from strain Lc 86 (Example 5). The partial sequence shows 61-68% sequence identity and 74-78% similarity with dextransucrases and alternansucrases (including dextransucrase DSRB742) of *Lc. mes.*

- 25 SEQ ID No's 11 and 12 give the nucleotide and amino acid sequence, respectively, of the glucansucrase of strain Lb180 (Example 2). The sequence shows 1322/1768 (74%) sequence identity and 1476/1768 (82%) similarity with 15/1768 gaps with glucansucrase from *Lb. reuteri* LB 121 as disclosed in WO 01/90372. The -35 and -10 sites TTGAAA and TATAA are located at nucleotide positions 561 and 599, respectively. The ribosome binding site (RBS) GAAGGAG is at 574 and the start codon ATG at 587. Inverted repeats AAGCAGCTC and GAGCTGCTT are at 6025 and 6051. Possible stop codons (TAA, TAG, TGA) are indicated with an * (5963).

- 30 SEQ ID No's 13 and 14 give the nucleotide and amino acid sequence, respectively, of the glucansucrase I from strain ML1 (Example 3). The sequence shows 1327/1775 (74%) sequence identity and 1465/1775 (81%) similarity with 17/1775 gaps with glucansucrase from *Lb. reuteri* LB 121 as disclosed in WO 01/90372, and 43-44% sequence identity and 57-58% similarity with dextransucrases of *Lc. mes.* and 47% sequence identity and 61% similarity with an alternansucrases of *Lc. mes.* The RBS AAGGAGA is at 31 and the start codon ATG is at 43. A stop codon TAG is at 5356.

35 SEQ ID No's 15 and 16 give the partial nucleotide and amino acid sequence, respectively, of a second glucansucrase from strain ML1 (ML4) (Example 3). The sequence shows

301/817 (36%) sequence identity and 427/817 (51%) similarity with 12/817 gaps with glucansucrase from *Lb. reuteri* LB 121 as disclosed in WO 01/90372, and 38% sequence identity and 53% similarity with glucosyltransferase of *Streptococcus mutans*.

5 SEQ ID No's 17 and 18 give the partial nucleotide and amino acid sequence, respectively, of the glucansucrase from strain LB 33 (Example 4). The sequence shows 59% sequence identity and 71% similarity with several known dextransucrases of *Lc. mes.* and 53% sequence identity and 67% similarity with other known dextransucrases (including dextransucrase DSRB742) of *Lc. mes.*

10 SEQ ID No's 19 and 20 give the nucleotide and amino acid sequence, respectively, of the glucansucrase from *Lb.* strain KG 15 (Example 6). The sequence shows 496/1111 (44%) sequence identity and 637/1111 (56%) similarity with 71/1111 gaps with glucansucrase from *Lb. reuteri* LB 121 as disclosed in WO 01/90372, and 57-59% sequence identity and 70% similarity with several dextransucrases (including dextransucrase DSRB742) of *Lc. mes.* The -35 and -10 sites *TTGGAC* and *TATTAT* are located at nucleotide positions 477 and 502, respectively. The RBS *GAAAGGA* is at 593 and the start codon *ATG* at 608. A stop codon *TAG* is 5393. Inverted repeats *AAAACAACCCCC* and *GGGGTTGTTTTT* are at 5497 and 5531 (-10.7 kcal/mole).

15 SEQ ID No's 21 and 22 give the partial nucleotide and amino acid sequence, respectively, of the glucansucrase from *Lb.* strain KG 3 (Example 7). The sequence shows 58 sequence identity and 71% similarity with known dextransucrases (including dextransucrase DSRB742) of *Lc. mes.*

Description of the figure

25 Figure 1 depicts an amino acid sequence alignment of glucosyltransferases (GTF) according to the invention. It shows the partial sequences of the GTF of *Lb* 180 (first line, starting with amino acid 216 of SEQ ID No. 12); GTF of ML1 (second line, starting with amino acid 15 of SEQ ID No. 14), GTF of *Lb* 33 (third line, starting with amino acid 222 or 243 of SEQ ID No. 18); GTF of KG15 (fourth line, starting with amino acid 567 of SEQ ID No. 20) and GTF of KG3 (fifth line, starting with amino acid 1 (LMAAF) of SEQ ID No. 22); and a GTF according to the invention of a *Lb. reuteri* strain "104" (sixth line, 1 (WPNTV) – 525). The alignment is not necessarily the best fit according to automated alignment programs, but is intended to define the enzymes of the invention.

35 The invention not only covers amino acid sequences shown in this figure, but also sequences wherein amino acids of a given sequence in the figure are exchanged with the corresponding amino acids (including gaps) of another sequence of the figure. This applies to stretches of at least 100 amino acids having at least 80%, preferably at least 90% identity with any of the sequences of the figure, or of the sequences listings given separately. It especially applies to the stretch of amino acids between the consensus peptides DNSN and YYGD (from 1202 to 1422 of SEQ ID No 12). Especially preferred

are sequences comprising the active core of the enzymes, which are present between the consensus peptides INGQ and VPDQ (from 957 to 1724 of SEQ ID No 12), with preferably at least 70% identity with any one of the core sequences given. A preferred non-identity with a given sequence is an exchange with the corresponding amino acids of another sequence. Especially preferred sequences are those where an amino acid at a given position is shared between at least 2, in particular at least 3, of the sequences of the figure. Most preferred are those sequences in which one of those consensus sequences is that of the GTF of Lb180, ML1 or Lb33 (first three lines). The N-terminal part upstream of the core (shown in the figure for GTF 180 and GTF ML1 only), or the C-terminal part downstream of the core (not shown in the figure) may be wholly or partly present or may be absent.

Claims

1. A process of producing a glucan having at least 10 anhydroglucose units, having a backbone consisting essentially of $\alpha(1,3)$ - and/or $\alpha(1,6)$ -linked anhydroglucose units (AGU), comprising subjecting sucrose to the activity of a glucosyltransferase produced by a *Lactobacillus* strain capable of producing $\alpha(1,3)$ - and/or $\alpha(1,6)$ -linked glucans, or to the *Lactobacillus* strain capable of expressing the glucosyltransferase.
2. A *Lactobacillus* strain capable of producing, in the presence of sucrose, a glucan having at least 10 anhydroglucose units (AGU) having a backbone consisting essentially of $\alpha(1,3)$ - and/or $\alpha(1,6)$ -linked AGU.
3. A glucan capable of being produced by glucosyltransferase activity of a lactic acid bacterium on a sucrose substrate, the glucan having an average molecular weight between 10 kDa and 1 GDa, especially between 10kDa and 50 MDa, and having a backbone consisting essentially of $\alpha(1,3)$ - and $\alpha(1,6)$ -linked anhydroglucose units (AGU).
4. A glucan according to claim 3, which is capable of being produced by glucosyltransferase activity of a *Lactobacillus* species.
5. A glucan according to claim 4, comprising 15-80% of $\alpha(1,3)$ -linked AGU, 2-80% of $\alpha(1,6)$ -linked AGU, and 2-25% of $\alpha(1,3,6)$ -linked AGU.
6. A glucan according to claim 5, having an average molecular weight of 50 kDa - 1 MDa and comprising 30-45% of $\alpha(1,3)$ -linked AGU, 30-45% of $\alpha(1,6)$ -linked AGU, and 3-13% of $\alpha(1,3,6)$ -linked AGU.
7. A glucan according to claim 5, having an average molecular weight of 10-50 MDa and comprising 15-26% $\alpha(1,3)$ -linked AGU, 30-50% of $\alpha(1,6)$ -linked AGU, 5-20% of $\alpha(1,3,6)$ -linked AGU.
8. A glucan according to claim 5, having an average molecular weight of 1-50 MDa and comprising 45-60% of $\alpha(1,3)$ -linked AGU, 4-10% of $\alpha(1,6)$ -linked AGU, and 10-20% of $\alpha(1,3,6)$ -linked AGU.

9. A glucan capable of being produced by glucosyltransferase activity of a lactic acid bacterium on a sucrose substrate, having an average molecular weight of 10-50 MDa and comprising 80-99% of $\alpha(1,6)$ -linked AGU and 0-15% of $\alpha(1,3)$ -linked AGU.
10. A protein having glucosyltransferase activity, capable of producing, in the presence of sucrose, a glucan according to any one of claims 3-9.
11. A protein according to claim 10, comprising an amino acid sequence of at least 100 amino acids exhibiting at least 70%, preferably at least 80%, amino acid identity with any one of the amino acid sequences of SEQ ID No. 2, 4, 8, 10, 12, 14, 16, 18, 20 and 22, and/or having a stretch of 100 amino acids having at least 80%, preferably at least 90%, amino acid identity with any one of the said amino acid sequences, or having at least 99% amino acid identity with the amino acid sequence of SEQ ID No. 6, and/or having a stretch of 100 amino acids having 100% amino acid identity with the amino acid sequence of SEQ ID No. 6.
12. A nucleic acid sequence encoding a protein according to claim 11.
13. A recombinant host cell containing one or more copies of a nucleic acid construct comprising a nucleic acid sequence according to claim 12 and capable of expressing a protein having glucosyl-transferase activity.
14. A *Lactobacillus* strain, capable of producing a glucan according to any one of claims 3-9, especially a *Lactobacillus* strain corresponding to strain 33, 180 or ML1 as described herein.
15. A *Leuconostoc* strain, capable of producing a glucan according to claim 9, especially a *Leuconostoc* strain corresponding to strain 86, deposited under accession number LMG P-20350.
16. A chemically modified glucan, which is obtained by 2,3-oxidation, 6-oxidation, phosphorylation, acylation, alkylation, hydroxyalkylation, carboxymethylation, amino-alkylation of one or more AGU of a glucan according to any one of claims 3-9.
17. Use of a glucan according to any one of claims 3-9, as a thickener.
18. Use of a glucan according to any one of claims 3-9, as a prebiotic and/or as a bioactive agent.

19. Use of a glucan according to any one of claims 3-9, as an anti-corrosion agent.
20. Use of a *Lactobacillus* bacterium capable of producing a glucan according to any one of claims 3-9, as a probiotic agent, or together with an indigestible glucan, as a synbiotic agent.

FIG. 1 SEQUENCE ALIGNMENT

```

216 MEIKKHFKLYKSGKQWVTAAVATVAVSTALLYGGVAHADQQVQSSTTQEQTSTVNADTTK
15  MEIKKHFKLYKSGKQWVTAAVATVAVSTALLYGGVAHADQQVQSSTTQDQTSTVNTNTTK

276 TVNLDTNTDQPAQTDDKNQVANDTTTNQSKTDSTSTTVKNPTFIPVSTLSSSDNEKQSQN
75  TIAADTNADQPAQTADKNQAASNDTTTNQSKTDSTSTTVKNLTSTPVSTLPSTDNEKQSQN

336 YNKPDNGNYGNVDAAYFNNNQLHISGWHATNASQGTDSRQVIVRDIITKTELGRTNVTNN
135 YNKHDNGNYGNIDTAYFSNNQLHVSQGNATNASQGTNSRQIIVRDIITNNELGRTDVTNN

396 VLRPDVKNVHNVYNADNSGFDVNINIDFSKMKDYRDSIEIVSRYSNGKSVDWWSQPITF
195 VARPDVKNVHNVYNADNSGFDINVNIEFSKMKDYRDSIEIVSRYSNGKSIDWWSQPITF

456 DKNNYAYLDTFEVKNGELHATGWNATNKAINYNHHFVILFDRGTNGKEVTRQEVDRDQGSRP
255 DKNNYAYLDTFEVKNGELHATGWNATNSAINYNHHFVILFDQTNGKEVARQEVREGQSRP

516 DVAKVYPQVVGANNSGFDVTFNIGDLDYTHQYQILSRYSNADNGEGDYVTYWFAPQSIAP
315 DVAKVYPQVVGADNSGFDVTFNIGNLDYTHQYQVLSRYSNSDNGEGDNVTYWFNPQSIAP

576 ANQSNQGYLDSFDISKNGEVTVTGWNATDLSELQTNHYVILFDQTAGQQVASAKVDLISR
375 ANQSNQGYLDSFDISKNGEVTVTGWNATDLSELQNNHYVILFDQTAGKQVASAKADLISR

636 PDVAKAYPTVKTAETSGFKVTFKVSNLQPGHQYSVVSFRFSADENGNGNDRHTDYWYSPV
435 PDVAKAYPTVKTAANSFGFKVTFKVNLDLQPGHQYSVVSFRFSADENGNGNDRHTDYWYSPV

696 TLNQATASNIDTITMTSNGLHITGWMASDNSINEATPYAIIILNNGREVTRQKLTLIARPDV
495 TLNQNASNIDTITMTSNGLHIGSWMASDNSINETTPYAIIILNNGKEVTRQKMSLTARPDV

756 AAVYPSLYNSAVSGFDTTIKLTNAQYQALNGQLQVLLRFSKAVDGNPNTVTVDQFSKN
555 AAVYPSLYNSAVSGFDTTIKLTNDQYQALNGQLQVLLRFSKAADGNPSGDNTVTVDQFSKN

816 YATTGGNFDYVKVNGNQIEFSGWHATNQSNQKDSQWIIIVLVNGKEVKRQLVNDTKDGAAG
615 YATTGGNFDYVKVNGNQVEFSGWHATNQSNQKDSQWIIIVLVNGKEVKRQLVNDTKEGAAG

876 FNRNDVYKVNPAIENSIMSGFQGIITLPVTVKDENVQLVHRFSNDAKTGEGNYVDFWSEV
675 FNRNDVYKVNPAIENSSMSGFQGIITLPVTVKENVQIVHRFSNDAKTGEGSHVDFWSEV

936 MSVKDSFQKGNGLNQFGLQTINGQYYIDPTTGQPRKNFLLQNGNDWIYFDKDTGAGTN
735 MPVKDSFQKGNGLKQFGLQTINGHQYYIDPMTGQPRKNFLLQNGNDWLYFDNETGEGTN
222 VNGKIYFVGDNQVKKNFTAIINGQSLYFNKTTGELASNDVQYENGLVKINDV
567 QTIAGKTYFFDKD GHLRKGYSTIIDNQLYYFDLKTGESVS

996 ALKLQFDKGTISADEQYRRGNEAYSYYDDKSIENVNGYLTADTWYRPKQILKDGTWTWDSK
795 ALKRQFDGGTISADSQYRKNEAGYDNKSIENVGGLTADTWYRPKQILKW TTWDSK
275 HNAAYSIDP?GFTNVNGFLTANSWYRPKYIYKDGQKWVEST
607 TTTSNFKSGLTSQTDDTTPHNSAVNMSKDSFTTVDGFLTAESWYVPKDIQTSATDWRAS

1056 ETDMPILMVWWPNTVTQAYYLNMYKQYGNLLPASLPSFSTADSAELNHYSELVQQNIE
854 ETDMPRLMVWWPNTVTQAYYLNMYKQHGNLLPANLPFFNSDADPLELNYYAEIVQQNIE
316 SQDMPRLMTWWPDKNTQVAYLQYM QKMILPADVTISSQTNQSVLTKESEF ITQAEIE
666 PEDFRPIMMTWWPTKQIQAAAYLNHVMSEG LLSSDKKFSATD DQTLNQAHAHVQLQIE
(0)
1 WPNTVTQAYYLNMYKQHGNLLPASLPPFNADADPAELNHYSEIVQQNIE

```

1116 KRISET GSTDWLRTLMEHFVTKNSMWNKDSENVYGGQLQGGFLKYVNSDLTKYANSW
914 KKISQT GNTDWLRTLMEHFVSNNMTWNKSENEDFGGLQLQGGFLKYVNSDKTPNANSW
374 KQIGVTNGNTDWLKKDISDFVNSQPNWNIDSEAKGTDH LQGGALLYVNNKLTPTYANSY
725 LKIQQT KSVEWLRTTMHNFIKSQPGYNVTSETPSNDH LQGGALSYINSVLTDPANSF
1 LMAAFVVTQPQWNKTSSEVDNDDH LQGGALTFENNGDT DANSY
50 KRISET GNTDWLRTLMEHFVTKNSMWNKDSENVYGGQLQGGFLKYVNSDLTKYANSW

1176 RLMNRTATNIDGKNY GGAEFLLANDIDNSNPVVQAEELNWLYYLMNFGTITGN
974 RIMGRQPANIDGNP IGSEFLLANDVDNSNPVVQAEQLNLWLYLLNFGTITAN
433 RLLNRTLNTQQGQVKDTS KQGGYEMLLANDVDNSNPVVQAEQLNLWLYYMMNIGSITAN
783 RLMNRNPTQQDGT RHYNTDTSEGGYELLLANDVDNSNPVVQAEQLNLWLYFLTHFGEIVKN
44 RLMNRTPTNQTGERLYHIDDSLGGYELLLANDVDNSNPQVQAEQLNLWLYYLMHFGDITAD
110 RLMDRATNIDGKNY GGAEFLLANDIDNSNPVVQAEELNWLYYLMNFGTITGN

1229 NPEANFDGIRVDAVDNVDVLLSIARDYFNAAYNMEQSDASANKHINILEDWGWDDPAYV
1027 DPANFDSIRVDAVDNVDADLLDIAGDYFNAVYHSQSNDKIANAHINILEDWGGQDPYYT
491 DPTANFDGYRVDVDNVDADLLNIAADYAKAYKTN QSDANANKHLSILEDWDDNDPAYI
843 DPSANFDSVRVDAVDNVDADLLNITAAYFRDVYGVKNDLTANQHLSILEDWGHNDPLYV
104 DPANFDAIRIDVDNVDADLLQLAAQYFRDAYGMATTDATSNKHLILEDWSHNDPAYM
163 NPEANFDGIRVDAVDNVDVLLSIARDYFNAAYNMEQSDANANKHINILEDWGWDDPAYV

1289 NKIGNPQLTMDRLRNAIMDTLSGAPDKNQALNKLITQSLVNRANDN TENAVIPSYNFV
1087 QSIGTPQLSMDYNFSTIRSVLASNTASMTD IKNSLVNRSLDN AENVSI PNYSFI
551 KAHGNNQLTMDFFPAHLAIKYSLNMPVSQSRGLEPELTTSLVNRTGDDSTENVAQPNYTFI
903 KDHGSDQLTMDDMHTQLIWSLTKNPDNRSAMRRFMEYYLVDRAKDN TSDPAIPNYSFV
164 QAHGNDQLTMDDMHTQLIWSLTKEAQRGTMARFMDFYLTNRANDD TENTAQPSYSFV
223 NKIGNPQLTMDRLRNAIMDTLSGAPDKNQALNKLITQSLVNRANDN TENAVIPSYNFV

1348 RAHDSNAQDQIRQAIQAATGKPYGE FNLDDEKKGMEAYINDQNSTNKKWNLYNMPSAY
1142 RAHDNGSQDDIKRAISDVNNLPYGSK FNFEQEQKGIEAYIADQSNVNKKWNLYNIPSSY
611 RAHDSEVQTIIAQIIKDKINPNSDGLTVPDEISQAFKIYNADLKTQYTFYNMPSAY
962 RAHDSEVQTVIGDIVAKLYPDVKNSL PSMEQLAAAFKVYDADMNSVNKKYTQYNMPAAY
223 RAHDSEVQTVIAEIVTKLHPEAGNGLMPTEEQMAEAFKIYNADQKKAVKTYTHYNMPSAY
282 RAHDSNAQDQIRQAIQAATGKPYGE FNLDDEKKGMEAYINDQNSTNKKWNLYNMPSAY

1406 TILLTNKDSVPRVYGGDLYQDGGQYMEHKTRYFDTITNLLKTRVKYVAGGQTMVSVDKN
1201 AIMLTNKDTVPRVYGGDLFTDGGQYMAQTTRYYPALTSLLKARIKYVAGGQTMVSVDKN
671 TILLTNKDTVPRVYGGDLYSDNGNYMSAHPYDAITLLKTRMKYVSGGQNMRMQYMQG
1021 AMLLTNKDTIPRVYGGDMYTDDGQYMATKSPYDAISALLKARIKYVAGGQTMVAVDKH
283 AMLLTNKDVIPIRYGGDLYTDDGQFMATKSPYFDAISTMLQARTKYVAGGQTMVAVDQH
340 TILLTNKDSVPRVYGGDLYQDGGQYMEHKTRYFDTITNLLKTRVKYVAGGQTMVSVDKN

1464 GILTNVRFKGAMNATDTGTDETRTEGIGVVISNNTNLKLNLDGESVVLHMG
1259 NILTSVRFKGAMNPTDMGDSLRTSGVGVIISNNDKLLSSNDKVVLHMG
731 DDMPANSYKGVLT SVRYGKGEMTADEQGNSETRTQGIGVII SNNPNLKLDSDNDQVVLNMG
1079 DILTSVRFGDGIMNASDKGSTTARTQGIGVIVSNNDALAL KGDTVTLHMG
341 DVLTSVRFKGAMTANDLGDAETRTGEGVLIISNNPKLQLGQQDNVVLHMG
398 GILTSVRFKGAMNATDTGTDETRTEGIGVVISNNTNLKLNLDGESVVLHMG

1515 AAHNQKYRAVILTTEGDKVKNYNTDTPAVAYTDANGDLHFTNTNLDG QQYTAVRG
1310 AAHNQKFKAVLLTTNDGIQSF NDDNAPVAYTDANGDLVLSGKDITTDGVIQHNTAVKG
791 AAHENQTYRPVLLTTKDGLKNYDSOSSVPQNALVSTNDKGQLIFKASS IQG
1129 IAHANQAYRALLTTTDLGIMKYTSDNGAPIRYTDANGDLIFTSADI KG
392 LAHANQAFRAVVLTTATGLTIY NDDDAPIRYTDNKGDLIFTNHDV YG
449 AAHNQKYRAVILTTEGDKVKNYNTDTPAVAYTDANGDLHFTNTNLDG QQYTAVRG

1571 YANPDVTGYLAVWVPAGAADDQDARTAPSDEAHTTKTAYRSNAALDSNVIYEGFSNFIYW
1369 YANADVKGYLAVWVPVGASVQQDIRTAPSGVQSDGKSVYHSNAALDSNII FEGFSNFVYW
842 VSNPQVSGYLSVWVPVGAKDNQDARTASSSQPSTDGKTYHSNAALDSQVIYEGFSNFQSI
1177 YQNVEVSGFLSVWVPVGASDTQDARATGSSAANKTGD^UTLHSNAALDSNVIYEGFSNFQEM
439 VLNPQVSGFLAMWVPTGAPANQDARSTASTNMSTDGSAYHSNAALDSQVIFESFSNFQAM
505 YANPDVTGYLAVWVPAGAADD

1631 PTTESERTNVRIAQNADLFKSWGITT^UFELAPQYNSSKDGTFLDSIIDNGYAFTDRYDLGM
1429 PTNNSERANVKIAQNTDLFKELGITSFELAPQYNSSKDGTFLDSQIDNGYAFTDRYDLGM
902 PTNTEFTNVKIAQNANLFKSLGITSFELAPQYRSSNDNSFLDSVVQNGYAFTDRYDIGY
1237 PTAHDEFTNVKIAQNADLFKSWGVT^USFQLAPQYRSSDDTSFLDSIIKNGYAFTDRYDLGF
499 PTSHDTYTNVVLANHADQLHDWGITSVQLAPQYRSSSTDGTFLDAIQNGYAFTDRYDLGF

1691 STPNKYGSDEDLRNALQALHKAGLQAIADWVPDQIYNLPGKEAVTVTRSDDHGTTWEVSP
1489 SIPNKYGS^UDTDLRNAIKALHKAGIQAMADWVPDQIYNLPGKEVVTATRVDERGNDWNVAQ
962 NTPTKYGTVTQLLDALRALHANGIQAI^UDDWVPDQIYNLPGEEIVAAQRTNGSGTYDQDSV
1297 NTPTKYGDVDDLADAI^URAMHSVGIQVMADFVPDQIYNLPGQEVVAVNRTNNFGTPNQDSD
559 GTPTKYGDDTDLRNVIKALHANGMQVMADFVPDQLYTLPGKELVQVTRTNMGEPT^UTHSD

SEQUENCE LISTING

<110> TNO

<120> Novel glucans and novel glucansucrases derived from
lactic acid bacteria

<130> Novel glucans and glucansucrases

<140>

<141>

<160> 10

<170> PatentIn Ver. 2.1

<210> 1

<211> 665

<212> DNA

<213> Lactobacillus reuteri

<400> 1

```

gataatacga atccggtggt gcaagctgaa gaattaaact ggctttacta tttaatgaat 60
ttcgggtacaa ttacaggaaa taatcctgaa gctaattttg atggtattcg agtggatgct 120
gttgataatg tagatgttga cttattgagt attgcacgtg attactttaa tgcagcatat 180
aacatggagc aaagtgatgc cagtgcctaat aagcacatta atattttgga agattgggga 240
tgggatgatc ctgcttatgt aaataagatt ggaaatcctc aattaacaat ggatgatcgt 300
ttacgaaatg caattatgga tacattatca ggagcacctg ataaaaacca agcattgaat 360
aaattaatta ctgagtcatt agtaaatcgt gctaattgata atactgaaaa cgcggttatt 420
ccaagctata attttggtcg agcacatgat agtaattgctc aagaccaa atcgtcaggct 480
attcaagctg caactggaaa accatatggc gaatttaact tagatgatga aaagaagggt 540
atggaagcat atattaatga tcagaattct actaataaga agtggaatct ttacaatatg 600
ccttctgctt atactattct tctaacaat aaagattcag ttccccacgt ctactatggc 660
gacat 665

```

<210> 2

<211> 221

<212> PRT

<213> Lactobacillus reuteri

<400> 2

```

Asp Asn Thr Asn Pro Val Val Gln Ala Glu Glu Leu Asn Trp Leu Tyr
 1                    5                    10                    15

```

```

Tyr Leu Met Asn Phe Gly Thr Ile Thr Gly Asn Asn Pro Glu Ala Asn
      20                    25                    30

```

```

Phe Asp Gly Ile Arg Val Asp Ala Val Asp Asn Val Asp Val Asp Leu
      35                    40                    45

```

```

Leu Ser Ile Ala Arg Asp Tyr Phe Asn Ala Ala Tyr Asn Met Glu Gln
      50                    55                    60

```

```

Ser Asp Ala Ser Ala Asn Lys His Ile Asn Ile Leu Glu Asp Trp Gly
      65                    70                    75                    80

```

```

Trp Asp Asp Pro Ala Tyr Val Asn Lys Ile Gly Asn Pro Gln Leu Thr
      85                    90                    95

```

```

Met Asp Asp Arg Leu Arg Asn Ala Ile Met Asp Thr Leu Ser Gly Ala

```


100	105	110
Pro Asp Lys Asn Gln Ala Leu Asn Lys Leu Ile Thr Gln Ser Leu Val		
115	120	125
Asn Arg Ala Asn Asp Asn Thr Glu Asn Ala Val Ile Pro Ser Tyr Asn		
130	135	140
Phe Val Arg Ala His Asp Ser Asn Ala Gln Asp Gln Ile Arg Gln Ala		
145	150	155
Ile Gln Ala Ala Thr Gly Lys Pro Tyr Gly Glu Phe Asn Leu Asp Asp		
165	170	175
Glu Lys Lys Gly Met Glu Ala Tyr Ile Asn Asp Gln Asn Ser Thr Asn		
180	185	190
Lys Lys Trp Asn Leu Tyr Asn Met Pro Ser Ala Tyr Thr Ile Leu Leu		
195	200	205
Thr Asn Lys Asp Ser Val Pro His Val Tyr Tyr Gly Asp		
210	215	220

<210> 3

<211> 674

<212> DNA

<213> Lactobacillus strain LB 33

<400> 3

```

gacaattcga atccggtggt gcaagcggaa cagttaaact gggtatacta catgatgaat 60
ataggtagca ttactgccaa tgatccacc gcaaactttg atggctatcg agtggacgct 120
gtggacaatg tcatgctga tttattaaat atagctgccg attatgccaa agatgcttat 180
aaaactaatc aaagtgatgc taatgccaac aaacatttat caatattaga agattgggat 240
aataatgac cggcttatat caaagcacat ggaaataatc agttaactat ggatttccca 300
gcacatttag caattaaata ttcattaaat atgccagtaa gtcaacgaag tgggctggaa 360
ccagagctca caaccagttt agttaacaga actggtgatg attctactga aaatgtcgca 420
cagccaaact atacttttat tagggctcac gatagtgaag tgcaaacaaat catcgcaaaa 480
attatcaaag ataaaatcaa ccctaactct gacggattaa cagttactcc cgatgaaata 540
agtcaggcct ttaaaatata taatgcagat gaattaaaga ctgataaaca atatactttt 600
tataacatgc cctctgccta tactattttg ctaaccaata aagatacagt accccacctc 660
tattacggcg acat                                     674

```

<210> 4

<211> 224

<212> PRT

<213> Lactobacillus strain LB 33

<400> 4

Asp Asn Ser Asn Pro Val Val Gln Ala Glu Gln Leu Asn Trp Leu Tyr		
1	5	10
Tyr Met Met Asn Ile Gly Ser Ile Thr Ala Asn Asp Pro Thr Ala Asn		
20	25	30
Phe Asp Gly Tyr Arg Val Asp Ala Val Asp Asn Val Asp Ala Asp Leu		
35	40	45

-3/28

50		55		60
Ser 65	Asp	Ala	Asn	Ala
		Asn	Lys	His
			Leu	Ser
			Ile	Leu
				Glu
				Asp
				Trp
				Asp
				80
Asn	Asn	Asp	Pro	Ala
				Tyr
				Ile
				Lys
				Ala
				His
				Gly
				Asn
				Asn
				Gln
				Leu
				Thr
				95
Met	Asp	Phe	Pro	Ala
				His
				Leu
				Ala
				Ile
				Lys
				Tyr
				Ser
				Leu
				Asn
				Met
				Pro
				110
Val	Ser	Gln	Arg	Ser
				Gly
				Leu
				Glu
				Pro
				Glu
				Leu
				Thr
				Thr
				Ser
				Leu
				Val
				125
Asn	Arg	Thr	Gly	Asp
				Asp
				Ser
				Thr
				Glu
				Asn
				Val
				Ala
				Gln
				Pro
				Asn
				Tyr
				140
Thr	Phe	Ile	Arg	Ala
				His
				Asp
				Ser
				Glu
				Val
				Gln
				Thr
				Ile
				Ile
				Ala
				Gln
				160
Ile	Ile	Lys	Asp	Lys
				Ile
				Asn
				Pro
				Asn
				Ser
				Asp
				Gly
				Leu
				Thr
				Val
				Thr
				175
Pro	Asp	Glu	Ile	Ser
				Gln
				Ala
				Phe
				Lys
				Ile
				Tyr
				Asn
				Ala
				Asp
				Glu
				Leu
				190
Lys	Thr	Asp	Lys	Gln
				Tyr
				Thr
				Phe
				Tyr
				Asn
				Met
				Pro
				Ser
				Ala
				Tyr
				Thr
				205
Ile	Leu	Leu	Thr	Asn
				Lys
				Asp
				Thr
				Val
				Pro
				His
				Leu
				Tyr
				Tyr
				Gly
				Asp
				220

<210> 5

<211> 671

<212> DNA

<213> Leuconostoc strain 86

<400> 5

```

gacaacacga acccggtggt gcaagctgag cagttaaatt ggttacacta tttgatgaat 60
tttggtagca ttacagcaaa cgattctgct gctaattttg atgagatacg tgtcgatgct 120
gtcgataatg ttgacgctga ttactccag attgcagcag attatttcaa agctgcttat 180
ggtgttgata aaaatgacgc aacagcaaat caacatcttt caattcttga agattggagc 240
cataacgacc ctgaatacgt gaaggatttt ggtaataatc aactcacaat ggatgattac 300
atgcataccc agttaatctg gtcgttgact aaagatatgc gtatgcgtgg taccatgcaa 360
cgcttcatgg actattacct cgtcaatcgc aatcacgata gtaccgaaaa cactgccatt 420
ccaaattaca gctttgttcg cgcacacgat agtgaagtac aaacagtcac tgcctcaaatt 480
atttctgagt tacatcccga cgtaaaaaat agtttggcac caacagcaga ccagctagcc 540
gaagccttta aagtttataa taacgatgaa aaacaggcgg ataagaaata tacacaatac 600
aacatgccta gcgcctatgc gatgctgtta actaataaag atacagtacc ccgcgtctac 660
tatggcgaca t

```

671

<210> 6

<211> 223

<212> PRT

<213> Leuconostoc strain 86

4/28

<400> 6

Asp Asn Thr Asn Pro Val Val Gln Ala Glu Gln Leu Asn Trp Leu His
 1 5 10 15
 Tyr Leu Met Asn Phe Gly Ser Ile Thr Ala Asn Asp Ser Ala Ala Asn
 20 25 30
 Phe Asp Glu Ile Arg Val Asp Ala Val Asp Asn Val Asp Ala Asp Leu
 35 40 45
 Leu Gln Ile Ala Ala Asp Tyr Phe Lys Ala Ala Tyr Gly Val Asp Lys
 50 55 60
 Asn Asp Ala Thr Ala Asn Gln His Leu Ser Ile Leu Glu Asp Trp Ser
 65 70 75 80
 His Asn Asp Pro Glu Tyr Val Lys Asp Phe Gly Asn Asn Gln Leu Thr
 85 90 95
 Met Asp Asp Tyr Met His Thr Gln Leu Ile Trp Ser Leu Thr Lys Asp
 100 105 110
 Met Arg Met Arg Gly Thr Met Gln Arg Phe Met Asp Tyr Tyr Leu Val
 115 120 125
 Asn Arg Asn His Asp Ser Thr Glu Asn Thr Ala Ile Pro Asn Tyr Ser
 130 135 140
 Phe Val Arg Ala His Asp Ser Glu Val Gln Thr Val Ile Ala Gln Ile
 145 150 155 160
 Ile Ser Glu Leu His Pro Asp Val Lys Asn Ser Leu Ala Pro Thr Ala
 165 170 175
 Asp Gln Leu Ala Glu Ala Phe Lys Val Tyr Asn Asn Asp Glu Lys Gln
 180 185 190
 Ala Asp Lys Lys Tyr Thr Gln Tyr Asn Met Pro Ser Ala Tyr Ala Met
 195 200 205
 Leu Leu Thr Asn Lys Asp Thr Val Pro Arg Val Tyr Tyr Gly Asp
 210 215 220

<210> 7

<211> 746

<212> DNA

<213> Leuconostoc strain 86

<400> 7

gactcactat agggcggaatt gggccctcta gatgcatgct cgagcggccg ccagtgtgat 60
 ggatatctgc agaattcgcc cttgacaatt ggaatccggt ggtgcaagct gaacagttga 120
 attggctaca ctatttgatg aatcttgga cgaattacagc taatgatcca gatgctaatt 180
 ttgacagcat aagagttgac gctgtcgata atgtggatgc agatttggtta gatattgcac 240
 gtgattactt taatgcagta tacaagggtta accaaagtga tgttaatgct aataaacata 300
 tttctatttt agaagattgg agtggattag atcccaatga ggttggttaa aatgggaatc 360
 cacaattaac acttaacaca ggggttcaaa attcattatt aaatgctttg acaaaggggc 420
 caaataatcg ttggggggat agactcattg attgataaat caacaatgaq atatccagat 480
 aaggatggta aaatccttat tcttaattat agtttcgtac gtgcacacga taqtgaaqtt 540

5/28

caaggtatta ttggcaaata ttaacagatc atacgtcagc cgaatcaggt aataaattca 600
 caaaggatcc attaaaacag gcttggatta ttactatgct gaatcaagaw tagactgtta 660
 aagaatattc gcactataat atggcgagtg cttatgcagc attgttaaca aattaagata 720
 ccattcccaa ctctactacg gcgact 746

<210> 8

<211> 221

<212> PRT

<213> Leuconostoc strain 86

<400> 8

Asp Asn Trp Asn Pro Val Val Gln Ala Glu Gln Leu Asn Trp Leu His
 1 5 10 15

Tyr Leu Met Asn Leu Gly Thr Ile Thr Ala Asn Asp Pro Asp Ala Asn
 20 25 30

Phe Asp Ser Ile Arg Val Asp Ala Val Asp Asn Val Asp Ala Asp Leu
 35 40 45

Leu Asp Ile Ala Arg Asp Tyr Phe Asn Ala Val Tyr Lys Val Asn Gln
 50 55 60

Ser Asp Val Asn Ala Asn Lys His Ile Ser Ile Leu Glu Asp Trp Ser
 65 70 75 80

Gly Leu Asp Pro Asn Glu Val Val Lys Asn Gly Asn Pro Gln Leu Thr
 85 90 95

Leu Asn Thr Gly Val Gln Asn Ser Leu Leu Asn Ala Leu Thr Lys Gly
 100 105 110

Pro Asn Asn Arg Trp Gly Ile Asp Ser Leu Ile Asp Lys Ser Thr Met
 115 120 125

Arg Tyr Pro Asp Lys Asp Gly Lys Ile Leu Ile Pro Asn Tyr Ser Phe
 130 135 140

Val Arg Ala His Asp Ser Glu Val Gln Gly Ile Ile Gly Lys Ile Leu
 145 150 155 160

Thr Asp His Thr Ser Ala Glu Ser Gly Asn Lys Phe Thr Lys Asp Gln
 165 170 175

Leu Lys Gln Ala Leu Asp Tyr Tyr Tyr Ala Asp Gln Asp Lys Thr Val
 180 185 190

Lys Glu Tyr Ser His Tyr Asn Met Ala Ser Ala Tyr Ala Ala Leu Leu
 195 200 205

Thr Asn Lys Asn Thr Ile Pro Asn Leu Tyr Tyr Gly Asp
 210 215 220

<210> 9

<211> 670

<212> DNA

<213> Leuconostoc strain 86

6/28

<400> 9

gataattcga atccgatggt gcaagctgag caattgaact ggcttcacta catgatgaac 60
 attggtacta tagctcagaa cgacccaaca gctaattttg acggttatcg tgttgatgcg 120
 gttgataatg ttgatgccga tctcttaciaa attgctgggtg attactttta agctgcatac 180
 ggtactggta aaactgaggc aaacgcaaac aatcatattt cgatcttgga agattgggat 240
 aataatgatt ctgcgtacat taaagcccac gggaacaacc aattgacaat ggattttcca 300
 gcacacttgg ctttgaaata cgccttgaac atgcctcttg ccgcacaaaag tggcctagaa 360
 ccgctaatta atacaagtct tgttaagcgt gggaaagatg ccacagaaaa tgaagcaciaa 420
 ccaaactatg cctttatccg tgcccatgat agtgaagtgc agacagttat tgcacaaatt 480
 attaaggata aaattaacac aaaatcagac ggcttaactg taacaccaga tgagattaag 540
 caagctttca atattttaciaa cgccgatgaa ttaaaagcag ataaggaata tacagcatac 600
 aatattcctg cttcttacgc tgtattgttg acaaacaagg atactgtgcc cgcgtctact 660
 atggcgacct 670

<210> 10

<211> 223

<212> PRT

<213> Leuconostoc strain 86

<400> 10

Asp Asn Ser Asn Pro Met Val Gln Ala Glu Gln Leu Asn Trp Leu His
 1 5 10 15
 Tyr Met Met Asn Ile Gly Thr Ile Ala Gln Asn Asp Pro Thr Ala Asn
 20 25 30
 Phe Asp Gly Tyr Arg Val Asp Ala Val Asp Asn Val Asp Ala Asp Leu
 35 40 45
 Leu Gln Ile Ala Gly Asp Tyr Phe Lys Ala Ala Tyr Gly Thr Gly Lys
 50 55 60
 Thr Glu Ala Asn Ala Asn Asn His Ile Ser Ile Leu Glu Asp Trp Asp
 65 70 75 80
 Asn Asn Asp Ser Ala Tyr Ile Lys Ala His Gly Asn Asn Gln Leu Thr
 85 90 95
 Met Asp Phe Pro Ala His Leu Ala Leu Lys Tyr Ala Leu Asn Met Pro
 100 105 110
 Leu Ala Ala Gln Ser Gly Leu Glu Pro Leu Ile Asn Thr Ser Leu Val
 115 120 125
 Lys Arg Gly Lys Asp Ala Thr Glu Asn Glu Ala Gln Pro Asn Tyr Ala
 130 135 140
 Phe Ile Arg Ala His Asp Ser Glu Val Gln Thr Val Ile Ala Gln Ile
 145 150 155 160
 Ile Lys Asp Lys Ile Asn Thr Lys Ser Asp Gly Leu Thr Val Thr Pro
 165 170 175
 Asp Glu Ile Lys Gln Ala Phe Asn Ile Tyr Asn Ala Asp Glu Leu Lys
 180 185 190
 Ala Asp Lys Glu Tyr Thr Ala Tyr Asn Ile Pro Ala Ser Tyr Ala Val
 195 200 205
 Leu Leu Thr Asn Lys Asp Thr Val Pro Ala Ser Thr Met Ala Thr

SEQ ID No. 11 DNA

SEQ ID No. 12 PRT

Lactobacillus reuteri strain 180

```
1   GAATTCGCTACCGAGATCTTAAAGGATAAAGACGTTGAAGTCGCTCCCGTTGATTCTTCA
1   N S L P R S * R I K T L K S L P L I L Q

61  GCATATCCAGCCAAAGCTTACCGTCCTCGCCATTCAAGTTATGAGTTTGAAGAAGGCAGAA
21  H I Q P K L T V L A I Q L * V * R R Q K

121 GACACTGGGTTTGAGATTATGGATTGGCGGACTGCATTGAGTAAGTTTATAGAGGGGATT
41  T L G L R L W I G G L H * V S L * R G L

181 GAGGAGTAAGATACTGGAACCGGTTTGGATTGGATACTGCTTTTTTATGGGCGGCGCAAT
61  R S K I L E P V W I G Y C F F M G G A I

241 AAAGCTAGATCTAACTGGAAAAGACTGCGAACAAAATTGAAATTTAGTGTAAGCAGCTAA
81  K L D L T G K D C E Q N * N L V * A A N

301 TATCCTTAGTCAATGTAGTATAATTGCAAATTTTTTACTAGGTAAGAAAGTATATTGTGG
101 I L S Q C S I I A N F L L G K K V Y C G

361 AAATATTTAAGAATATTGTCGTTACCGGTAGAGACAATTTTATAAGTTCTAACTTTGTTC
121 N I * E Y C R Y R * R Q F Y K F * L C S

421 ACTATGTTGTTAACCCTTACTAGGAAGTTGAACATATTACGGTTTTAGATAAGTTAACTT
141 L C C * P L L G S * T Y Y G F R * V N L

481 ATACTGGCATTTAGTCAATTCTGATATCTTTGTTTAAAATTACAAATTTGAACTTTGTTT
161 Y W H L V N S D I F V * N Y K F E L C L
      - 35

541 GAAGAAAATGTGGGAAGAATTTGAAAATTTCTTTTAAAAAATTAAACATCATAGTATTA
181 K K M W E E F E N F L * K N * T S * Y Y
      -10                               RBS      start

601 TAATATCGATAATTAAATTGTTTATTCTGACATGAAGGAGATTAAATGGAAATAAGAA
201 N I D N * I V Y S D M K E I K M E I K K

661 ACATTTTAAAGTTGTATAAAAGTGGTAAACAATGGGTGACAGCGGCAGTTGCTACTGTTGC
221 H F K L Y K S G K Q W V T A A V A T V A

721 CGTTTCAACCGCGCTTCTTTACGGGGGAGTTGCGCATGCTGATCAACAAGTTCAGTCCTC
241 V S T A L L Y G G V A H A D Q Q V Q S S

781 CACAACCCAAGAACAACTTCTACTGTGAATGCTGATACTACTAAACAGTAAATTTAGA
261 T T Q E Q T S T V N A D T T K T V N L D

841 TACTAATACTGACCAACCAGCCCAAACAACCTGATAAAAATCAAGTAGCAAATGACACTAC
281 T N T D Q P A Q T T D K N Q V A N D T T

901 TACTAACCAAAGTAAACTGATAGTACATCAACAACCTGTTAAGAATCCTACTTTTATACC
301 T N Q S K T D S T S T T V K N P T F I P

961 AGTTTCTACTTTGTCTTCATCAGATAATGAAAAACAAAGTCAAAATTATAATAAGCCGGA
321 V S T L S S S D N E K Q S Q N Y N K P D

1021 TAATGGAACTATGGAAATGTTGATGCAGCTTACTTTAATAATAATCAATTGCATATTTTC
341 N G N Y G N V D A A Y F N N N Q L H I S

1081 AGGATGGCACGCAACAAATGCATCTCAAGGAACAGATAGTCGTCAGGTGATTGTACGTGA
361 G W H A T N A S Q G T D S R Q V I V R D
```

1141 TATCACAACTAAAACCTGAATTAGGACGTACTAATGTAACAAACAATGTTTTACGCCCAGA
381 I T T K T E L G R T N V T N N V L R P D

1201 TGTAA[~]AAATGTCCACAATGTTTATAACGCTGATAATTCTGGATTTCGATGTCAACATCAA
401 V K N V H N V Y N A D N S G F D V N I N

1261 CATTGACTTTAGTAAGATGAAGGACTATCGTGATTCAATTGAAATTGTTAGTCGATACAG
421 I D F S K M K D Y R D S I E I V S R Y S

1321 TGGAAATGGTAAATCTGTTGATTGGTGGTCTCAACCGATTACCTTTGACAAAAATAATTA
441 G N G K S V D W W S Q P I T F D K N N Y

1381 CGCATACCTTGACACATTTGAAGTTAAAAATGGGGAATTGCATGCAACAGGATGGAATGC
461 A Y L D T F E V K N G E L H A T G W N A

1441 TACTAATAAGGCAATTAACATAACCACCATTTTGTAAATTTTATTTGATCGAACAAATGG
481 T N K A I N Y N H H F V I L F D R T N G

1501 TAAAGAAGTGACTCGTCAAGAAGTTCGTGATGGTCAATCGCGTCCAGATGTTGCTAAGGT
501 K E V T R Q E V R D G Q S R P D V A K V

1561 ATATCCACAAGTAGTTGGGGCAAATAACTCTGGCTTTGACGTGACATTTAATATTGGTGA
521 Y P Q V V G A N N S G F D V T F N I G D

1621 TCTAGATTACACTCATCAATACCAAATTCCTTAGTCGTTACAGCAATGCAGATAATGGCGA
541 L D Y T H Q Y Q I L S R Y S N A D N G E

1681 AGGTGATTATGTTACTTACTGGTTTGCTCCACAATCAATTGCTCCTGCTAACCAAAGTAA
561 G D Y V T Y W F A P Q S I A P A N Q S N

1741 TCAGGGTTATTTAGATTTCATTTGATATTAGTAAAAATGGTGAAGTGACAGTAACTGGTTG
581 Q G Y L D S F D I S K N G E V T V T G W

1801 GAATGCTACTGATCTATCTGAATTACAACTAACCATTATGTAATTTTATTTGACCAAAC
601 N A T D L S E L Q T N H Y V I L F D Q T

1861 CGCTGGTCAACAAGTTGCATCTGCAAAAGTTGATCTAATTTCCCGTCCAGATGTTGCGAA
621 A G Q Q V A S A K V D L I S R P D V A K

1921 AGCTTACCCAACAGTAAAAACTGCTGAACTTCTGGCTTTAAGGTAACATTTAAGGTTAG
641 A Y P T V K T A E T S G F K V T F K V S

1981 TAATTTACAACCAGGTCATCAATATAGTGTCGTAAGCCGTTTTTCTGCCGATGAAAACGG
661 N L Q P G H Q Y S V V S R F S A D E N G

2041 TAATGGTAATGATAAACGTCATACCGATTACTGGTACAGCCCAGTAACCTTAAATCAAAC
681 N G N D K R H T D Y W Y S P V T L N Q T

2101 TGCTTCAAATATTGATACTATCACAATGACATCGAATGGATTGCATATTACTGGTTGGAT
701 A S N I D T I T M T S N G L H I T G W M

2161 GGCAAGTGATAATTCAATTAATGAAGCAACTCCATATGCCATTATTCTTAATAATGGTAG
721 A S D N S I N E A T P Y A I I L N N G R

2221 AGAGGTTACTCGTCAAAAATTAAC[~]TTAATTGCGCGTCCAGATGTAGCAGCAGTATATCC
741 E V T R Q K L T L I A R P D V A A V Y P

2281 TTCACTCTATAACAGTGCTGTTAGTGGATTTGATACTACCATTAAGTTGACTAATGCTCA
761 S L Y N S A V S G F D T T I K L T N A Q

2341 ATACCAGGCGCTTAATGGTCAACTACAAGTATTGTTACGTTTTTCTAAAGCTGTTGATGG
781 Y Q A L N G Q L Q V L L R F S K A V D G

9/28

2401 TAATCCAAACGGCACTAATACTGTAACAGATCAATTTAGTAAGAATTATGCAACTACTGG
801 N P N G T N T V T D Q F S K N Y A T T G

2461 TGGAAACTTTGATTATGTCAAAGTAAACGGCAATCAAATTGAATTTAGTGGCTGGCATGC
821 G N F D Y V K V N G N Q I E F S G W H A

2521 AACTAATCAATCAAATGATAAAAAATTCTCAATGGATTATTGTTTTAGTTAATGGTAAAGA
841 T N Q S N D K N S Q W I I V L V N G K E

2581 GGTAAAACGGCAATTAGTTAATGATACTAAGGATGGTGCTGCTGGGTTCACCGTAATGA
861 V K R Q L V N D T K D G A A G F N R N D

2641 TGTTTACAAAGTAAATCCGGCTATTGAAAATAGTATTATGTCTGGGTTCGAAGGTATTAT
881 V Y K V N P A I E N S I M S G F Q G I I

2701 TACTTTACCTGTAACAGTTAAGGATGAAAATGTTTCAGCTTGTTTCATCGTTTTAGTAATGA
901 T L P V T V K D E N V Q L V H R F S N D

2761 TGCAAAGACTGGTGAAGGTAATTATGTTGATTTCTGGTCAGAAGTAATGTCTGTTAAGGA
921 A K T G E G N Y V D F W S E V M S V K D

2821 CAGCTTCCAAAAGGGTAATGGTCCGCTTAATCAATTTGGTTTACAACTATTAACGGCCA
941 S F Q K G N G P L N Q F G L Q T I N G Q

2881 ACAATATTATATTGACCCAACAACCTGGCCAACCTCGTAAGAATTTCTTATTGCAAAATGG
961 Q Y Y I D P T T G Q P R K N F L L Q N G

2941 GAACGATTGGATTTACTTTGACAAAGATACTGGTGCTGGAACATAATGCTCTTAAGTTACA
981 N D W I Y F D K D T G A G T N A L K L Q

3001 ATTTGATAAGGGAACAATTTCTGCTGATGAGCAATATCGTCGAGGAAATGAAGCCTATAG
1001 F D K G T I S A D E Q Y R R G N E A Y S

3061 TTATGATGACAAGAGTATTGAAAATGTAAATGGTTACTTAAACAGCTGATACTTGGTACCG
1021 Y D D K S I E N V N G Y L T A D T W Y R

3121 ACCAAAACAAATCTTAAAGGATGGTACTACTTGGACTGACTCTAAAGAAACAGATATGCG
1041 P K Q I L K D G T T W T D S K E T D M R

3181 CCCAATTTTAAATGGTATGGTGGCCAAATACTGTTACACAAGCATATTATCTTAACTACAT
1061 P I L M V W W P N T V T Q A Y Y L N Y M

3241 GAAGCAATATGGTAATTTATTGCCGGCTAGTTTACCAAGCTTCAGTACAGATGCTGATTC
1081 K Q Y G N L L P A S L P S F S T D A D S

3301 TGCTGAATTAAATCATTACTCCGAGCTTGTTCAACAAAATATCGAAAAGCGGATCAGTGA
1101 A E L N H Y S E L V Q Q N I E K R I S E

3361 GACTGGTAGTACTGATTGGTTACGTACACTAATGCATGAGTTCGTTACTAAGAATTCTAT
1121 T G S T D W L R T L M H E F V T K N S M

3421 GTGGAATAAGGATAGTGAAAATGTGCGATTACGGTGGTTTGCAATTACAAGGTGGATTCCT
1141 W N K D S E N V D Y G G L Q L Q G G F L

3481 TAAGTATGTAAATAGTGATCTTACTAAATATGCAAATTCAGATTGGCGTTTAAATGAACCG
1161 K Y V N S D L T K Y A N S D W R L M N R

3541 TACAGCTACTAATATTGATGGTAAGAACTATGGTGGTGCGGAATTCTTATTAGCTAATGA
1181 T A T N I D G K N Y G G A E F L L A N D

3601 TATTGATAACTCAAATCCAGTTGTTCAAGCTGAAGAATTAACTGGCTTTACTATTTAAT
1201 I D N S N P V V Q A E E L N W L Y Y L M

10/28

3661 GAATTTGGGTACAATTACAGGAAATAATCCTGAAGCTAATTTTGATGGTATTTCGAGTGGA
1221 N F G T I T G N N P E A N F D G I R V D

3721 TGCTGTTGATAATGTAGATGTTGACTTATTGAGTATTGCACGTGATTACTTTAATGCAGC
1241 A V D N V D V D L L S I A R D Y F N A A

3781 ATATAACATGGAGCAAAGTGATGCCAGTGCTAATAAGCACATTAATATTTTGAAGATTG
1261 Y N M E Q S D A S A N K H I N I L E D W

3841 GGGATGGGATGATCCTGCTTATGTAAATAAGATTGGAAATCCTCAATTAACAATGGATGA
1281 G W D D P A Y V N K I G N P Q L T M D D

3901 TCGTTTACGAAATGCAATTATGGATACATTATCAGGAGCACCTGATAAAAAACCAAGCATT
1301 R L R N A I M D T L S G A P D K N Q A L

3961 GAATAAATTAATTACTCAGTCATTAGTAAATCGTGCTAATGATAATACTGAAAACGCGGT
1321 N K L I T Q S L V N R A N D N T E N A V

4021 TATTCCAAGCTATAATTTTGTTCGAGCACATGATAGTAATGCTCAAGACCAAATTCGTCA
1341 I P S Y N F V R A H D S N A Q D Q I R Q

4081 GGCTATTCAAGCTGCAACTGGAAAAACCATATGGCGAATTTAACTTAGATGATGAAAAGAA
1361 A I Q A A T G K P Y G E F N L D D E K K

4141 GGGTATGGAAGCATATATTAATGATCAGAATTCTACTAATAAGAAGTGAATCTTTACAA
1381 G M E A Y I N D Q N S T N K K W N L Y N

4201 TATGCCTTCTGCTTATACTATTCTTCTAACAAATAAAGATTTCAGTTCCTCGTGTTTACTA
1401 M P S A Y T I L L T N K D S V P R V Y Y

4261 TGGAGACCTCTACCAAGATGGTGGTCAATATATGGAACATAAAACACGTTACTTTGATAC
1421 G D L Y Q D G G Q Y M E H K T R Y F D T

4321 TATTACTAACTTATTAAAGACACGGGTTAAATATGTTGCCGGTGGACAAACTATGAGTGT
1441 I T N L L K T R V K Y V A G G Q T M S V

4381 TGATAAGAATGGTATTCTTACAAACGTTTCGTTTTGGGAAAGGCGCCATGAATGCTACTGA
1461 D K N G I L T N V R F G K G A M N A T D

4441 TACTGGTACTGATGAAACAAGAACAGAAGGTATCGGTGTTGTAATTAGTAACAATACTAA
1481 T G T D E T R T E G I G V V I S N N T N

4501 TTTGAAGCTTAATGATGGTGAATCAGTAGTGCTTCATATGGGAGCTGCTCATAAGAATCA
1501 L K L N D G E S V V L H M G A A H K N Q

4561 AAAGTATCGTGCTGTGATCTTAACAACCTGAAGATGGTGTGAAGAATTACACTAATGATAC
1521 K Y R A V I L T T E D G V K N Y T N D T

4621 AGACGCACCAGTTGCATACACTGATGCTAATGGTGACCTTCACCTTACTAATACTAATTT
1541 D A P V A Y T D A N G D L H F T N T N L

4681 AGATGGTCAACAATATACAGCTGTTTCGTGGATATGCAAATCCTGATGTAACAGGATATCT
1561 D G Q Q Y T A V R G Y A N P D V T G Y L

4741 AGCTGTTTGGGTACCAGCTGGAGCAGCAGATGATCAAGATGCACGTACTGCACCAAGTGA
1581 A V W V P A G A A D D Q D A R T A P S D

4801 TGAGGCCCATACTACAAAGACTGCTTATCGCTCTAATGCAGCCCTTGATTCTAACGTTAT
1601 E A H T T K T A Y R S N A A L D S N V I

4861 TTATGAAGGATTCTCTAACTTCATTTACTGGCCAACCTACTGAAAGCGAACGGACTAATGT
1621 Y E G F S N F I Y W P T T E S E R T N V

11/28

4921 GAGAATTGCACAAATGCGGATCTATTTAAGTCATGGGGAATTACTACCTTTGAATTAGC
1641 R I A Q N A D L F K S W G I T T F E L A

4981 TCCACAATACAATTCAAGTAAAGATGGTACGTTCTTGATTCAATAATTGATAATGGATA
1661 P Q Y N S S K D G T F L D S I I D N G Y

5041 TGCCTTTACTGATCGTTATGATTTAGGAATGAGTACTCCTAACAAGTATGGATCTGATGA
1681 A F T D R Y D L G M S T P N K Y G S D E

5101 AGACTTACGTAATGCTTTACAAGCCTTACATAAAGCTGGTTTACAAGCAATTGCCGACTG
1701 D L R N A L Q A L H K A G L Q A I A D W

5161 GGTTCCTGATCAAATTTATAACTTACCTGGTAAAGAAGCTGTAACAGTAACACGTTTCTAGA
1721 V P D Q I Y N L P G K E A V T V T R S D

5221 TGATCACGGTACTACATGGGAAGTTTCGCCAATAAAGAATGTTGTCTATATTACAAATAC
1741 D H G T T W E V S P I K N V V Y I T N T

5281 GATTGGTGGAGGTGAATACCAGAAGAAATATGGTGGTGAATTCTTAGACACTCTTCAAAA
1761 I G G G E Y Q K K Y G G E F L D T L Q K

5341 AGAATATCCACAATTATTTAGTCAGGTATATCCAGTAACTCAAACGACAATTGATCCTAG
1781 E Y P Q L F S Q V Y P V T Q T T I D P S

5401 TGTTAAGATTAAAGAGTGGTCTGCTAAATACTTTAATGGTACTAATATCCTTCATCGAGG
1801 V K I K E W S A K Y F N G T N I L H R G

5461 TGCTGGATATGTATTGCGCTCTAATGATGGTAAATACTATAATCTTGGTACAAGCACTCA
1821 A G Y V L R S N D G K Y Y N L G T S T Q

5521 ACAATTCTTACCGTCTCAATTATCAGTTCAAGATAATGAAGGATATGGATTTGTAAAAGA
1841 Q F L P S Q L S V Q D N E G Y G F V K E

5581 AGGAAATAATTACCATTACTATGATGAGAATAAACAGATGGTAAAAGATGCGTTTATTCA
1861 G N N Y H Y Y D E N K Q M V K D A F I Q

5641 AGATAGTGTGGTAATTGGTATTACTTCGATAAAAAATGGTAATATGGTTGCTAACCAAAG
1881 D S V G N W Y Y F D K N G N M V A N Q S

5701 TCCTGTTGAAATTAGTAGTAATGGAGCTTCAGGAACCTACCTTTTCTTGAACAATGGGAC
1901 P V E I S S N G A S G T Y L F L N N G T

5761 ATCATTCCGTTCTGGATTGGTGAAACTGATGCAGGTACGTACTATTATGATGGCGATGG
1921 S F R S G L V K T D A G T Y Y Y D G D G

5821 CCGAATGGTTCGTAATCAAACGGTAAGTGATGGTGCGATGACATATGTTCTTGATGAAAA
1941 R M V R N Q T V S D G A M T Y V L D E N

5881 TGGTAACTTGTTAGTGAATCATTTGATTCTGCTACTGAAGCACACCCATTAAAACC
1961 G K L V S E S F D S S A T E A H P L K P

5941 TGGTGATTTAAACGGCCAAAAATAATTACAATATGAAAATTGGAACCTTGATTTTACCTT
1981 G D L N G Q ' K * L Q Y E N W N L Y F T F
inverted repeat

6001 CTTTGAAATAATATAGTTCTAATTAAGCAGCTCGCACCAAGACTTGGTATGAGCTGCTTT
2001 F E I I * F * L S S S H Q D L V * A A F

6061 TTTTGGCTCTACAATATCTGGTGTGATATAGAAATATCACTTTCTATACCAATATCAGA
2021 F G S T I S G V D I E I S L S I P I S D

6121 TTTTTGTTTTTAACTAAAAAGAGGCTCGCCCTCTGATACAATGAAATCGCCAAATCAC
2041 F C F * T K K E A R P L I Q * N R Q I T

12/28

6181 ATAGTAAAGAAGGTAACCTCCATGGATAATGATACAAGAAGCTCTTCTCAATTTAACAGAC
2061 * * R R * P P W I M I Q E L F S I * Q T

6241 CCTCATTTAAATTTTCCTCATCATTGGCTTAAATATAAAACAATTAAAAAAGTTCCGGTG
2081 L I * I F L I I G L N I K Q L K K F G W

6301 GCACAAATATNCTGTACCCTTTCTTATACACCACGGGCTTGTCCAAATTGGGGGAGTCA
2101 H K Y X V P F L I H H G X C P N W G S H

6361 TTAATCGNGGTCAAATCTTAAATATGGGCTTTTATCAAGCTAAACACAATATGGACAAT
2121 * S X S N L K I W A F I K L N T I W T I

6421 TTAAAACTCAACCATTAATGNTG
2141 * N S T I N X

SEQ ID No. 13 DNA

SEQ ID No. 14 PRT

Lactobacillus reuteri strain ML1

1 ATCGATAATCAAATTGTTTATTTTGATATAAAGGAGATTAAATGGAAATAAAGAAACAT
1 I D N Q I V Y F D I K E I K M E I K K H
RBS start

61 TTTAAGTTGTATAAAAGTGGTAAACAATGGGTGACAGCGGCTGTTGCTACTGTTGCCGTT
21 F K L Y K S G K Q W V T A A V A T V A V

121 TCAACCGCGCTTCTTTACGGGGGAGTTGCACATGCTGATCAACAAGTTCAGTCTTCCACA
41 S T A L L Y G G V A H A D Q Q V Q S S T

181 ACTCAAGACCAAACCTTCTACTGTAAATACTAATACTACTAAAACAATAGCTGCAGATACT
61 T Q D Q T S T V N T N T T K T I A A D T

241 AATGCTGATCAGCCAGCTCAAACAGCTGATAAAATCAAGCAGCATCAAATGACACTACT
81 N A D Q P A Q T A D K N Q A A S N D T T

301 AACCAAAGTAAACTGATAGTACTTCAACAACCTGTTAAGAATCTTACTTCTACACCAGTT
101 N Q S K T D S T S T T V K N L T S T P V

361 TCTACTTTGCCATCAACTGATAATGAAAAACAAATCAAATTTATAATAAGCATGATAAT
121 S T L P S T D N E K Q N Q N Y N K H D N

421 GGAAACTATGGGAATATTGATACTGCTTACTTTAGCAATAATCAATTGCATGTTTCAGGA
141 G N Y G N I D T A Y F S N N Q L H V S G

481 TGGAATGCAACGAATGCATCTCAAGGAACAAACAGTCGGCAAATTATTGTGCGTGATATC
161 W N A T N A S Q G T N S R Q I I V R D I

541 ACAACCAATAATGAATTAGGTCGTACTGATGTAACAAACAATGTTGCGCGCCCAGACGTT
181 T T N N E L G R T D V T N N V A R P D V

601 AAGAATGTTTCATAATGTTTATAACGCTGATAATTCTGGATTTGATATTAATGTCAATATT
201 K N V H N V Y N A D N S G F D I N V N I

661 GAATTTAGCAAGATGAAAGATTATCGGGATTCAATTGAAATTGTTAGTCGATACAGTGGA
221 E F S K M K D Y R D S I E I V S R Y S G

721 AACGGTAAATCTATTGACTGGTGGTCCCAACCGATCACTTTTGACAAAAACAATTATGCT
241 N G K S I D W W S Q P I T F D K N N Y A

781 TATCTTGATACATTTGAAGTGAAAAATGGCGAATTACATGCAACCGGATGGAATGCTACT
261 Y L D T F E V K N G E L H A T G W N A T

841 AATAGTGCAATTAACATAATCACCATTTTGTAAATTTTATTTGATCAAACGAATGGTAAG
281 N S A I N Y N H H F V I L F D Q T N G K

901 GAAGTAGCACGACAAGAAGTTCGTGAAGGCCAATCACGCCCAGATGTTGCTAAGGTATAT
301 E V A R Q E V R E G Q S R P D V A K V Y

961 CCACAAGTAGTTGGTGCTGACAACCTCCGGCTTTGATGTGACATTTAATATCGGTAATTTA
321 P Q V V G A D N S G F D V T F N I G N L

1021 GATTATACTCACCAGTACCAAGTTCTTAGTCGTTACAGCAATTCTGATAATGGCGAAGGC
341 D Y T H Q Y Q V L S R Y S N S D N G E G

1081 GATAATGTTACCTACTGGTTTAAATCCACAATCCATTGCTCCTGCTAATCAAAGTAACCAG
361 D N V T Y W F N P Q S I A P A N Q S N Q

1141 GGTATCTAGACTCATTTGATATTAGTAAAAATGGTGAAGTAACAGTGACCGGATGGAAT
381 G Y L D S F D I S K N G E V T V T G W N

1201 GCTACTGACTTGTGAGAATTACAAAATAACCATTATGTAATTCTATTTGATCAGACAGCA
401 A T D L S E L Q N N H Y V I L F D Q T A

1261 GGCAAACAAGTAGCATCTGCCAAGGCTGATTTAATTTACGTCCAGATGTTGCAAAGGCT
421 G K Q V A S A K A D L I S R P D V A K A

1321 TATCCAACAGTAAAACTGCTGCAAATCTGGCTTTAAGGTAACATTTAAGGTTAATGAT
441 Y P T V K T A A N S G F K V T F K V N D

1381 TTACAACCGGGTCACCAATATAGCGTTGTAAGTCGTTTCTCTGCCGATGAAAATGGTAAT
461 L Q P G H Q Y S V V S R F S A D E N G N

1441 GGTAATGATAAGCGTCATACAGATTACTGGTTTAGTCCAGTAACATTAAACCAGAATGCT
481 G N D K R H T D Y W F S P V T L N Q N A

1501 TCAAACATTGATACTATTACAATGACATCTAATGGGTTACATATTGGCAGTTGGATGGCA
501 S N I D T I T M T S N G L H I G S W M A

1561 AGTGATAACTCAATTAATGAAACAACCTCCATATGCTATTATTCTCAATAACGGTAAAGAA
521 S D N S I N E T T P Y A I I L N N G K E

1621 GTTACTCGTCAAAAGATGAGTTTAACTGCCCCGTCCAGATGTAGCAGCAGTATATCCTTCA
541 V T R Q K M S L T A R P D V A A V Y P S

1681 CTTTATAATAGTGCTGTTAGTGGGTTTGATACTACTATTAAATTGACTAATGATCAGTAT
561 L Y N S A V S G F D T T I K L T N D Q Y

1741 CAAGCGCTTAATGGTCAATTACAAGTATTGTTACGTTTTTCTAAAGCTGCTGATGGTAAT
581 Q A L N G Q L Q V L L R F S K A A D G N

1801 CCAAGTGGTGATAATACTGTAACCTGATCAATTTAGTAAAAATTATGCAACTACTGGTGGA
601 P S G D N T V T D Q F S K N Y A T T G G

1861 AACTTTGATTATGTAAAAGTAAATGGTAATCAAGTTGAATTTAGTGGTTGGCATGCAACT
621 N F D Y V K V N G N Q V E F S G W H A T

1921 AACCAATCAAATGATAAAGATTCACAATGGATTATTGTTTTAGTTAATGGTAAAGAAGTA
641 N Q S N D K D S Q W I I V L V N G K E V

1981 AAGCGTCAATTAGTTAATGATACTAAAGAGGGGGCTGCTGGCTTCAACCGAAACGATGTC
661 K R Q L V N D T K E G A A G F N R N D V

2041 TACAAAGTAAATCCAGCTATTGAAAACAGTTCTATGTCTGGATTCCAAGGCATTATTACT
681 Y K V N P A I E N S S M S G F Q G I I T

2101 TTACCAGTAACAGTTAAGAATGAGAATGTTTCAGATTGTCCATCGTTTTAGTAATGATGCA
701 L P V T V K N E N V Q I V H R F S N D A

2161 AAGACAGGTGAAGGTAGCCATGTTGATTTCTGGTCAGAAGTAATGCCAGTTAAGGATAGT
721 K T G E G S H V D F W S E V M P V K D S

2221 TTCCAAAAGGGTAATGGTCCGCTTAAGCAATTGGCTTACAACTATTAATGGTCATCAA
741 F Q K G N G P L K Q F G L Q T I N G H Q

2281 TATTATATTGACCCAATGACTGGCCAACCTCGCAAGAACTTCCTATTACAAAATGGTAAT
761 Y Y I D P M T G Q P R K N F L L Q N G N

2341 GACTGGCTTTATTTTGATAATGAACTGGTGAGGGAACATAATGCGTTAAAGAGGCAATTT
781 D W L Y F D N E T G E G T N A L K R Q F

2401 GACGGAGGAACGATTTCTGCTGATAGTCAGTATAGAAAGGGTAATGAAGCTTATGGTTAT
801 D G G T I S A D S Q Y R K G N E A Y G Y

2461 GACAATAAGAGCATTGAAAATGTTGATGGCTTTTTAACAGCTGATACTTGGTACCGACCA
821 D N K S I E N V D G F L T A D T W Y R P

2521 AAACAAATTTTAAAATGGACCACCTGGACAGATTCTAAAGAAACAGATATGCGACCGCTC
841 K Q I L K W T T W T D S K E T D M R P L

2581 TTAATGGTTTTGGTGGCCAAATACTGTAACCCAAGCATATTACCTTAACATACATGAAACAA
861 L M V W W P N T V T Q A Y Y L N Y M K Q

2641 CATGGAACTTATTACCAGCTAATCTTCCATTCTTTAATTCTGATGCAGATCCATTAGAA
881 H G N L L P A N L P F F N S D A D P L E

2701 TTAAATTATTATGCAGAAATTGTTTCAGCAAAATATTGAAAAGAAGATTAGTCAAACCTGGT
901 L N Y Y A E I V Q Q N I E K K I S Q T G

2761 AATACTGACTGGTTGCGAACTTTGATGCACGAATTTGTATCTAATAATACAATGTGGAAT
921 N T D W L R T L M H E F V S N N T M W N

2821 AAGAATAGTGAAAATGAAGACTTTGGTGGGTTGCAATTACAAGGTGGTTTTCTAAAGTAC
941 K N S E N E D F G G L Q L Q G G F L K Y

2881 GTTAATAGTGATAAGACACCTAATGCTAATTCTAATTGGCGTATTATGGGTAGGCAGCCA
961 V N S D K T P N A N S N W R I M G R Q P

2941 GCTAATATTGACGGAAATGGGCCAATTGGATCAGAATTCTTATTAGCTAATGACGTTGAT
981 A N I D G N G P I G S E F L L A N D V D

3001 AATTCTAATCCAGTTGTTCAAGCTGAACAGTTAAATTGGCTACATTACTTATTGAATTTT
1001 N S N P V V Q A E Q L N W L H Y L L N F

3061 GGAACTATTACTGCAAATGATCCTGATGCTAATTTTGATAGCATTTCGTGTTGATGCTGTT
1021 G T I T A N D P D A N F D S I R V D A V

3121 GACAATGTAGATGCCGATTTATTAGATATAGCTGGTGATTACTTTAATGCAGTATATCAT
1041 D N V D A D L L D I A G D Y F N A V Y H

3181 TCTCAAAGTAATGATAAAATTGCTAATGCTCATATTAATATTCTTGAGGATTGGGGTGGC
1061 S Q S N D K I A N A H I N I L E D W G G

3241 CAAGATCCGTATTATACGCAAAGCATCGGAACTCCTCAATTATCGATGGATTATAATTTTC
1081 Q D P Y Y T Q S I G T P Q L S M D Y N F

3301 TCAACTATAAGAAGTGTGTTAGCATCTAACACTGCATCAATGACTGATATTATTAAGAAT
1101 S T I R S V L A S N T A S M T D I I K N

3361 TCATTGGTAAATCGGAGCTTAGATAATGCTGAAAACGTATCAATTCCTAATTACTCATTT
1121 S L V N R S L D N A E N V S I P N Y S F

3421 ATCCGTGCACATGATAATGGTTCACAAGATGATATTAAGCGTGCAATTCAGATGTAAAT
1141 I R A H D N G S Q D D I R A I S D V N

3481 AATTTACCATATGGTTCGAAGTTTAACTTTGAGCAAAGCAAAAGGGGATTGAAGCATAC
1161 N L P Y G S K F N F E Q E Q K G I E A Y

3541 ATTCAGATCAAAGTAATGTTAATAAGAAGTGGGAATAATTATAATATTCCATCTTCATAT
1181 I A D Q S N V N K K W N N Y N I P S S Y

3601 GCTATTATGTTGACTAATAAGGATACCGTTCCTCGTGTATATTATGGTGATTATTTACT
1201 A I M L T N K D T V P R V Y Y G D L F T

3661 GATGGTGGTCAGTATATGGCACAACAACGCGTTATTATCCTGCACCTTACAAGTCTTTTA
1221 D G G Q Y M A Q T T R Y Y P A L T S L L

3721 AAGGCACGTATTAAGTATGTAGCTGGTGGACAAACAATGTCTGTGCGATAAGAATAATATT
1241 K A R I K Y V A G G Q T M S V D K N N I

3781 TTGACTAGTGTTCGCTTTGGTAAAGGTGCGATGAATCCTACTGATATGGGTGATAGTTTA
1261 L T S V R F G K G A M N P T D M G D S L

3841 ACTAGAACATCTGGTGTGGGGTAGTTATAAGTAATAATGATAAATTATTATTAAAGCTCA
1281 T R T S G V G V V I S N N D K L L L S S

3901 AATGATAAAGTTGTATTACACATGGGTGCTGCACATAAGAATCAGAAATTTAAAGCAGTC
1301 N D K V V L H M G A A H K N Q K F K A V

3961 TTACTAACTACTAATGATGGTATTCAGAGTTTAAATGATGACAATGCGCCTGTTGCATAT
1321 L L T T N D G I Q S F N D D N A P V A Y

4021 ACTGATGCTAATGGTGACTTGGTCCTTTCTGGTAAAGATATTACGACTGATGGTGTAATT
1341 T D A N G D L V L S G K D I T T D G V I

4081 CAACATAATACTGCTGTTAAGGGCTATGCTAATGCTGATGTTAAAGGTTATCTTGCAGTA
1361 Q H N T A V K G Y A N A D V K G Y L A V

4141 TGGGTTCCAGTAGGTGCCAGTGTACAACAGGATATTAGAACAGCACCATCAGGGGTACAA
1381 W V P V G A S V Q Q D I R T A P S G V Q

4201 AGTGATGGAAAGTCTGTTTATCATTCAAATGCAGCTCTGGATTCAAATATTATTTTGAA
1401 S D G K S V Y H S N A A L D S N I I F E

4261 GGATTCTCTAACTTTGTATATTGGCCGACAAATAATTCTGAGCGTGCAAATGTAAAAATC
1421 G F S N F V Y W P T N N S E R A N V K I

4321 GCTCAGAATACTGACTTATTTAAGGAGTTGGGTATTACTTCATTTGAATTAGCTCCACAG
1441 A Q N T D L F K E L G I T S F E L A P Q

4381 TATAATTCAAGTAAGGATGGCACATTCCTTGATTCTCAGATTGATAATGGATATGCATT
1461 Y N S S K D G T F L D S Q I D N G Y A F

4441 ACTGATCGCTATGATCTAGGTATGAGCATTCCAAATAAGTATGGTAGCGATACTGATCTA
1481 T D R Y D L G M S I P N K Y G S D T D L

4501 AGGAATGCTATTAAAGCCTTACATAAGGCCGAATTCAAGCAATGGCTGATTGGGTTTCCT
1501 R N A I K A L H K A G I Q A M A D W V P

4561 GATCAAATTTATAATTTACCAGGTAAAGAAGTTGTTACTGCTACTCGTGTGACGAACGT
1521 D Q I Y N L P G K E V V T A T R V D E R

4621 GGAAATGATTGGAATGTAGCTCAGATTAAGGATTCACCTTTATGTTGCTAATACAATTGGT
1541 G N D W N V A Q I K D S L Y V A N T I G

4681 GGTGGAAAGTATCAAGAGCAATATGGTGGAGCTTTCCTTGATCAATTACAAAAGCAATAT
1561 G G K Y Q E Q Y G G A F L D Q L Q K Q Y

4741 CCACAAATCTTTGAACGTAAACAACCTTCAACTGGTGTAGCAATTGACCCAAGTACTAAG
1581 P Q I F E R K Q P S T G V A I D P S T K

4801 ATTAAACAGTGGTCTGCTAAATACTTTAATGGGACAAATATTTTACATCGTGGTGCAGGG
1601 I K Q W S A K Y F N G T N I L H R G A G

4861 TATGTATTAAGAGATAACGGTGGTAACTACTTTAGCCTTGGAATAGTAATAATAAACAG
1621 Y V L R D N G G N Y F S L G N S N N K Q

4921 TTATTACCAAATCAATTATCAGGTAAGGCTGAAAATGGCTTTGTTGATGTTAATGGGAAT
1641 L L P N Q L S G K A E N G F V D V N G N

4981 ACTAAATACTTTACATCAACCGGAATTCCTGTCACGGATGCATTGTTC AAGACAGTGTA
1661 T K Y F T S T G I P V T D A F V Q D S V

5041 GGTAAGTGGTACTATATTGATAAAAATGGTAATATGCTTAAAAATACCGGTTTTGTAGAT
1681 G N W Y Y I D K N G N M L K N T G F V D

5101 ATTACGCGAAATGGTCAGACAGGTACGTATCTATTCTTAAATAACGGTATCTCATTCCGA
1701 I T R N G Q T G T Y L F L N N G I S F R

5161 TCAGGATTAGTTAAAATTGGTAATGATACTTATTACTTTGACGGTAATGGAAAAATGGTT
1721 S G L V K I G N D T Y Y F D G N G K M V

5221 CGTGGCCAATCTATTAGTGATGGTACGATGAATTATACTCTTGATAAGGATGGTAAATTA
1741 R G Q S I S D G T M N Y T L D K D G K L

5281 GTTGGCTTGTATTATGATCCAAGTAGTCAGAATCCACATCCAATTACTCAACAGGATTTA
1761 V G L Y Y D P S S Q N P H P I T Q Q D L

5341 AGTGGTACTAATAAGTAGTTTATTAAAAATCACCAATAGAAGTTGTCTCTACATCAAATG
1781 S G T N K * F I K N H Q * K L S L H Q M

5401 GTGTTGATATGAAAATATAATACTTTTATACCATTAAATTGGTCTAGTAAGAATCATCCTC
1801 V L I * K Y N T L Y H * I G L V R I I L

5461 ACGGATGGTTCTTTTTAGTTTCGCCGTTTGTAATAAAGTTAGAAAAATAAAAAAGCCA
1821 T D G S F * F R R L * N * V R K N K K P

5521 TTTGTGATAGACTTTTGAGTATCCCTAATCAAAAGAAAGGCAATCACAAATGACCTATAA
1841 F V I D F * V S L I K R K A I T N D L *

5581 ACATCTTACCACACGCGAATTAACCTCTCATAGCTGATTTTTGGTATCAAGGCACTAAAGC
1861 T S Y H T R I N S H S * F L V S R H * S

5641 TTATCGGGCTGCTAAATTACTTCAACGTAGTCAAGAAACCATCTATCGTGTTTATCGTTT
1881 L S G C * I T S T * S R N H L S C L S F

5701 CCTCAATAACGGTAAAACCATCGACCAATATCTTCAGACTTATCAGCGACATAAACGTCG
1901 P Q * R * N H R P I S S D L S A T * T S

5761 TTGTGGTTCGGAAGCAGACCCAAGTCCCACTATCGAGGTTAACTATATCCATGCGCAAAT
1921 L W S E A D P T A N Y R G * L Y P C A N

5821 CAAGGCTGGTTGGACTCCTGATACTATTATTGGTCGTGATGAGCACCCGATTAGCTGCAG
1941 Q G W L D S * Y Y Y W S * * A P D * L Q

5881 ATACTAATGCTGATCAGCCAGCTCAAACAGCTGATAAAAATCAAGCAGCATCAAATGACA
1961 I L M L I S Q L K Q L I K I K Q H Q M T

5941 CTACTAACCAAAGTAAACTGATAGTACTTCAACAACTGGTAAGAATCTTACTTCTACAC
1981 L L T K V K L I V L Q Q L V R I L L L H

6001 CAGTTTTCTACTTTGGCATCAACTGATAATGGAAAACAAAATCAAAATTATAATAAGCAT
2001 Q F S T L A S T D N G K Q N Q N Y N K H

6061 GATAT
2021 D

SEQ ID No. 15 DNA

SEQ ID No. 16 PRT

Lactobacillus reuteri strain ML1 (ML4)

1 AATATTGATGGTTACTTAAGTTATACTGGTTGGTATCGTCCTTATGGAACGAGTCAAGAT
1 N I D G Y L S Y T G W Y R P Y G T S Q D

61 GGTAAACATGGTACGAAACAACGCAATGGATTGGCGTCCATTACTGATGTATATTTGG
21 G K T W Y E T T A M D W R P L L M Y I W

121 CCAAGCAAAGATGTTCAAGCACAATTTATTAAGTATTTTGTTAATAATGGTTATGAGAAT
41 P S K D V Q A Q F I K Y F V N N G Y E N

181 GCTAATTATGGACTTACAGAGTCCTCTGTTGCTTCCTTTAGCAAGGATACTAATGCTAAT
61 A N Y G L T E S S V A S F S K D T N A N

241 CTCCTCGATGTAACTGCACAAAATCTTCGTTATGTAATTGAGCAAAGTATTGCAGCCAAT
81 L L D V T A Q N L R Y V I E Q S I A A N

301 AAAGGGACAAGTAAGTTAGCAAATGATATTAATAGTTTTGCTGCAACGGTTCCTGAATTA
101 K G T S K L A N D I N S F A A T V P E L

361 TCTGCATCATCTGAATTATCATTGCAAAGCATGCCAAACTATCGACCAGATGAAAGTGGA
121 S A S S E L S L Q S M P N Y R P D E S G

421 ACTGTTGATAGTGATCAAGTCATTTTTGTTAATAATAATTCAAAGGATCCCCGTAAAGGG
141 T V D S D Q V I F V N N N S K D P R K G

481 AACACTGGTTATGCGGACAGCAACTATCGCTTAATGAACAGGACGATTAATAATCAGGCC
161 N T G Y A D S N Y R L M N R T I N N Q A

541 GGAAATAATAATAGTGATAACAGTCCAGAACTCCTTGTTGGTAATGATATTGATAATTCA
181 G N N N S D N S P E L L V G N D I D N S

601 AACCCAGTAGTACAAGCTGAAAATCTTAATTGGGAATACTTTTTACTAAATTATGGTAAG
201 N P V V Q A E N L N W E Y F L L N Y G K

661 TTAATGGGGTATAATCCAGACGGTAATTTTGATGGCTTCCGAGTTGATGCTGCTGATAAT
221 L M G Y N P D G N F D G F R V D A A D N

721 ATTGATGCAGATGTCTTAGATCAAATGGGTCAATTAATGAACGACATGTATCATACAAAG
241 I D A D V L D Q M G Q L M N D M Y H T K

781 GGAAATCCTCAAAATGCAAATGATCATTTGAGTTATAATGAAGGTTATCATTCTGGGGCT
261 G N P Q N A N D H L S Y N E G Y H S G A

841 GCACAAATGCTAAATGAAAAGGGTAATCCTCAATTGTACATGGATTCAGGCGAATTCTAT
281 A Q M L N E K G N P Q L Y M D S G E F Y

901 ACCCTTGAGAATGTTCTCGGACGTGCAAATAACCGTGATAGTATCGGTAATTTAATTACT
301 T L E N V L G R A N N R D S I G N L I T

961 AATAGTGTGTTAATCGGCAAAATGATACAACAGAGAATGAAGCTACGCCAAACTGGTCA
321 N S V V N R Q N D T T E N E A T P N W S

1021 TTTGTA[~]ACTAACCATGATCAACGAAAGAATTTGATTAATAGATTAATTATTAAGGGTCAT
341 F V T N H D Q R K N L I N R L I I K G H

1081 CCTAACATTCCGGATATTATGGGTTTACGCTTACAAAGCTGAATATGCAAATCAAGCATGG
361 P N I P D I M G S A Y K A E Y A N Q A W

1141 CAAGAATTCTACGCTGATCAGAAAAAGACTAATAACAATATGATCAATATAATGTTCCG
381 Q E F Y A D Q K K T N K Q Y D Q Y N V P

1201 GCTCAGTATGCAATTCTTTTGAGCAATAAAGATACGGTTCCGCAGGTTTACTATGGTGAC
401 A Q Y A I L L S N K D T V P Q V Y Y G D

1261 CTTTATAATGAAACTGCTCAATACATGCAAGAGAAGTCAATTTACTATGATACAATCACG
421 L Y N E T A Q Y M Q E K S I Y Y D T I T

1321 ACTCTTATGAAGGCCCGTAAACAATTTGTTAGTGGTGGTCAAACGATGACTAAACTTAAC
441 T L M K A R K Q F V S G G Q T M T K L N

1381 AATAATTTATTAGCTAGTGTTCGATATGGTAAGGGTGTGCTGATTCTAATAGCAATGGT
461 N N L L A S V R Y G K G V A D S N S N G

1441 ACCGATAAGCTTAGCCGAACAAGTGGGATAGCCGCTTAGTTGGTAATGATAGTAATATG
481 T D K L S R T S G I A V L V G N D S N M

1501 GCTCAACAAACTGTTGCTATTAATATGGGTCGCGCTCATGCTAACCAACAATATCGAAAT
501 A Q Q T V A I N M G R A H A N Q Q Y R N

1561 TTAATTGATACTACCGAAAATGGCTTGACATATGATGGAGAAAATAGTGAAAATCCAGCC
521 L I D T T E N G L T Y D G E N S E N P A

1621 ATTTTGACAACTGATAGTAATGGTATCTTAAAAGTAACAGTTAAAGGATACAGTAACCCA
541 I L T T D S N G I L K V T V K G Y S N P

1681 TACGTAAGTGGTTATCTTGGTGTGTTGGGTTCCAGTAATTTCTGGTGATCAAGATGTTACT
561 Y V S G Y L G V W V P V I S G D Q D V T

1741 ACAAGTGCAAGTGATGTTGTGCTGATAAAGAAAAGACTTTTGAATCTAATGCTGCTCTT
581 T S A S D V V A D K E K T F E S N A A L

1801 GATTCTCATATGATCTATGAAGATTTACGCTTGTTCCAACCAGAACCAACTAATGTTGAG
601 D S H M I Y E D F S L F Q P E P T N V E

1861 AATCATGCTTACAATGTGATTGCTAAAAATGCTAATCTCTTTAATGATTTAGGCATTACT
621 N H A Y N V I A K N A N L F N D L G I T

1921 GATTTTTGGATGGCTCCTGCTTACACTCCATTTGGAATGAGTCGTTATAATGAAGGATAC
641 D F W M A P A Y T P F G M S R Y N E G Y

1981 TCAATGACGGATCGTTACAATTTAGGTACGACAGCTAATCCAACAAAGTATGGTAGTGGA
661 S M T D R Y N L G T T A N P T K Y G S G

2041 GAAGAGCTTGCAAATACAATTGCTGCATTGCATAAAGTAGGATTAAAAGTTCAAGAAGAT
681 E E L A N T I A A L H K V G L K V Q E D

2101 ATTGTTATGAATCAGATGATTGGTTTCTCTGGTCAAGAAGCAGTAACGGTTACTCGAACA
701 I V M N Q M I G F S G Q E A V T V T R T

2161 AATAATCGTGAATGCAGATTCATGTAAATGGTCAAACATATGCAAATCAAATTTACTTT
721 N N R G M Q I H V N G Q T Y A N Q I Y F

19/28

2221 GCATATACAACCTGGTGGCGGAAATGGTCAAGAACTTATGGTGGTAAATACCTTGCCGAA
741 A Y T T G G G N G Q E T Y G G K Y L A E

2281 TTACAAAAGAACTATCCTGACCTATTTACGACCAAGGCAATTTGACAGAAGTTGTACCT
761 L Q K N Y P D L F T T K A I S T E V V P

2341 GATCCAACCGTTCGTATTAAT
781 D P T V R I N

SEQ ID No. 17 DNA
Lactobacillus strain LB33

1 ATGGAATTAA AAAGGCATTA CAAGATGTAC AAGGCTGGTA AAAAATGGGT TTTTGCTGCA
61 ATTGCCACAA TCTCTATAAT TGCAGGATTA AATACAGTGG CAGTGACAAC CTATGCTGCC
121 GGCAATAATG ATCCGCAGCA GACCACTACT CAAAATGCAC CTAACAACAG TAACGATCCG
181 CAATCTACTA CTACGCAGAA TACTGCCAAC AACAGTAACG ATCCGCAATC TACTACTACG
241 CAGAATACTG CCAACAACAG TAATGGTCCA CAATCTACTA CTACGCAGAA TACTGCCAAC
301 AATAGTAATG GTCCACAATC TACTACTACG CAGAATACTG CCAATAACAG TAACGATCCA
361 CAATCTACTA CTACGCAGAA TACTGCCAAC AACAGTAACG ATCCGCAATC TACTACTACG
421 CAGAATACTG CCAACAATAG TAATGGTCCA CAATCTACTA CTACGCAGAA TACTGCCAAC
481 AACAGTAACG ATCCGCAATC TACTACTACG CAAAACACTG CCAACAACGG TAATGATCCA
541 CAATCTACTA CTGGAAGAAGA TACAGTTAGT ATTGCAGATA TTCAAGTTAA CCAACCTGTT
601 AATCTTTTAG GAAAGCAATC AACTGTATCT AGTACTGGTT ATAATGACTC TCACATAAAA
661 AATGTCAATG GGAAAATCTA TTTTGTGGT GATAATGGTC AGGTCAGAA AAACTTTACA
721 GCCATAATCA ATGGACAATC ACTATATTTT AATAAAACAA CTGGAGAATT GGCTTCTAAT
781 GATGTTCAAT ATGAAAATGG GTTAGTAAAA ATAAACGATG TTCATAACGC CGCTTACTCT
841 ATTGATCCAA CGGGATTCAC TAATGTAAAC GGATTTTTAA CTGCTAATAG TTGGTATAGA
901 CCCAAATATA TTTACAAAGA TGGGCAAAAA TGGGTGGAAT CAACCTCTCA AGATATGCGT
961 CCCCTTTTAA TGACATGGTG GCCAGATAAA AATACTCAAG TAGCTTATTT ACAATATATG
1021 CAGAAAATGG GCATTTTACC CGCTGACGTC ACTATATCAA GTCAAACCAA TCAATCAGTT
1081 TTAACCAAAG AATCATTTAT TACTCAAGCT GAAATTGAAA AACAGATTGG AGTAAACAAAT
1141 GGAAACACTG ATTGGCTAAA GAAAGATATC TCTGATTTTG TAAATTCTCA ACCAAATTGG
1201 AATATAGATA GTGAAGCCAA AGGCACAGAC CATTTGCAGG GGGGAGCACT TTTATATGTT
1261 AATAATAAGT TAACTCCATA TCGGAATTC TATTACCGCT TGCTTAACCG AACACTTACT
1321 AATCAACAGG GGCAAGTAAA AGATACTTCT AAACAAGGCG GTTATGAAAT GTTACTTGCC
1381 AACGATGTGG ATAATTCCAA TCCAGTAGTT CAAGCGGAAC AGTTAACTG GTTATACTAC
1441 ATGATGAATA TAGGTAGCAT TACTGCCAAT GATCCCACCG CAAACTTTGA TGGCTATCGA
1501 GTGGACGCTG TGGACAATGT CGATGCTGAT TTATTAAATA TAGCTGCCGA TTATGCCAAA
1561 GATGCTTATA AAACATAATCA AAGTGATGCT AATGCCAACA AACATTTATC AATATTAGAA
1621 GATTGGGATA ATAATGATCC GGCTTATATC AAAGCACATG GAAATAATCA GTTAACTATG
1681 GATTTCCCAG CACATTTAGC AATTAAATAT TCATTAAATA TGCCAGTAAG TCAACGAAGT
1741 GGGCTGGAAC CAGAGCTCAC AACCAGTTTA GTTAACAGAA CTGGTGATGA TTCTACTGAA
1801 AATGTCGCAC AGCCAAACTA TACTTTTATT AGGGCTCACG ATAGTGAAGT GCAAACAATC
1861 ATCGCACAAA TTATCAAAGA TAAAATCAAC CCTAACTCTG ACGGATTAAC AGTTACTCCC
1921 GATGAAATAA GTCAGGCCTT TAAAATATAT AATGCAGATG AATTAAGAC TGATAAACAA
1981 TATACTTTTT ATAACATGCC CTCTGCCTAT ACTATTTTGC TAACCAATAA AGATACAGTA
2041 CCTCGAGTTT ATTATGGGGA TCTTTATAGT GATAATGGCA ATTATATGTC AGCCCATTCT
2101 CCTTACTATG ATGCAATAAC TACGTTATTA AAAACACGAA TGAAATACGT ATCTGGTGGT
2161 CAAAACATGC GTATGCAATA TATGCAGGGT GATGATATGC CTGCTAATAG CTATAAGGGC
2221 GTTTTAACTT CAGTTAGATA TGGTAAGGGT GAAATGACAG CCGATGAGCA AGGTAATTCA
2281 GAAACTCGTA CTCAAGGAAT TGGGGTCATT ATAAGCAATA ATCCTAATTT AAAATTAGAC
2341 AGTAATGACC AAGTGGTATT AAATATGGGG GCGGCACATG AAAATCAAAC TTATCGCCCT
2401 GTATTACTAA CAACTAAAGA TGGATTGAAA AACTATGATT CCGATAGTTC TGTACCTCAA
2461 AATGCATTAG TTTCAACCAA CGATAAGGGA CAACTCATAT TTAAAGCTAG TTCTATTAG
2521 GGAGTAAGTA ATCCGCAGGT ATCTGGTTAT TTGTCCGTGT GGGTCCCAGT GGGGGCAAAG
2581 GATAATCAAG ATGCTCGGAC TGCAAGCAGT TCTCAGCCAT CAACTGATGG GAAAACATAT
2641 CATTTCCAATG CTGCTTTAGA CTCTCAAGTT ATTTACGAAG GATTTTCTAA TTTTCAATCG
2701 ATTCCTACAA ATACAGAAGA TTTCACTAAT GTAAAAATTG CTCAAAACGC TAACCTGTTT
2761 AAGAGCTTGG GAATAACAAG TTTTGAATTA GCCCCTCAAT ATCGTTCCAG TAATGATAAT
2821 AGTTTTCTGG ATTCGGTTGT TCAAAATGGC TACGCATTTA CTGATCGTTA TGATATTGGG
2881 TATAATACTC CGACAAAATA TGGAACTGTT ACTCAATTGC TGGATGCATT AAGGGCTTTA
2941 CATGCCAACG GAATTCAGC GATCGATGAC TGGGTTCCCTG ACCAAATATA CAATTTACCT
3001 GGTGAGGAAA TTGTCGCAGC TCAAAGAACT AATGGATCTG GGACATATGA TCAAGATTCT

20/28

3061 GTTATTGATG ATACATTATA TGATTCTCAC ACTGTTGGTG GTGGCGAATA TCAAGCTAAA
 3121 TTTGGTGGAG CTTTTCTAAA CAAGTTAAAG CAGTTGTATC CTGATTTATT TAAAGTTAAA
 3181 CAAATTTCTA CTGGTCAACC TATGAATCCT AATGAAAGAA TTACCGAGTG GTCAGCAAAG
 3241 TACTTTAATG GTACAAATAT TCAAGGAAGA GGCGCTTGGT ATGTATTAAA AGACTGGGGT
 3301 ACCAATCAGT ACTTTAATGT AAGTAATAAC CAGTTTGTTT CCAAACAATT CCTAGGTACA
 3361 GATACTTATA CAGGCTTTAA TGTTACAAAT GAGGGAAGTC AGTTTTATTC TACGAGTGGG
 3421 TATAAAGCCC AGAATACCTT TATTCAGGAC GGAGACAACT GGTATTACTT TGACAATAAT
 3481 GGCTATATGG TAACTGGTTT ACAGAATATA AATGGGAATA ATTACTATTT CTGCCCCAAT
 3541 GGCATTGAAC TACAAGACTC TTATTTATTTG AATGATGATA CCGGTAAAGA ATATTATTAT
 3601 GCAAGTAATG GTAAGCAAAT CTCAAATCGT TATTATCCAG ATGCTAACCG CAATTGGAGA
 3661 TATTTCTTCA ATGATGGTTC AATGGCAAGA AATGGATTAA CCACTATTGA ACAACCAGAT
 3721 GGGCAAAAAG TGATCCAATA TTTTGATTCC GATGGTATTC AATTAAAGGG AAATGCCGCA
 3781 AAAGATAATA ATGGTAATTT AAGATATTTT GACGGTAATA CAGGTGATAT GGTCAATTAAT
 3841 TCATTTGGAG AACTTCCTGA TGGCTCTTGG TTATACCTTA ATGATAAGGG GATTGCCGTT
 3901 ACTGGTAAAC AGGAAATCAA TGGTCAAAACC TACTACTTTG ATGCGGATGG CAAGCAAGTG
 3961 AAGAATGATT TTAGAGAGTT GCCTGATGGT TCATGGCTTT ATCTTAATGA CAAGGGGATT
 4021 GCCGTTACTG GTAAACAGGA AATCAATGGT CAAACCTACT ACTTTGATGC GGATGGCAAG
 4081 CAACTGAAGA ATGATTTTAG AGAGTTGCCT GATGGTTCAT GGCTTTATCT TAATGACAAG
 4141 GGGATTGCCG TTAAGGTAA ACAGGGAATC AATGGTCAAA CCTATGCAGA GGCTAAAAATC
 4201 ACAGCTGCCG AAAATGCTCA TCAAGCTGCC ACAGACGCTG TGAATAAAGC CCAAGTGCCT
 4261 CAATCGCCTA ACACTAGTTC CTCTAGTTCT AGCGTTAGCC AAGCTACTAA ACATCAATTG
 4321 GCAGTTAAAA CTGCTAAAGC TCAACTTGCT AAAACTAAGG CTCAAAATTGC TAAGTATCAA
 4381 AAGGCTTTGA AAAAAAGCCAA AACTACAAAG GCCAAGGCTC AAGCTCGTAA AAGTTTGAAG
 4441 AAGGCCGAGA CTAGTTTCAG CAAAGCTGAA CTTAATTTGG CATTATTAAA TAATAAAGCC
 4501 GTAAAAGCTG CACAACTAA GGTTAATAAG GCTAAGGCTC AAGTCACTAA ATACCAAAAG
 4561 GCTTTGAAGA AAGCTAAGAC TACAAAGGCT AAGACTCAAG CTCGTAAAAA TTTGAAGAAG
 4621 GCCAACTCTA GTCTGACAAA AGCTCAAAAA GCATTAACTA AAGTAATTAA AACCAATATC
 4681 AAGTAA

SEQ ID No. 18 PRT

Lactobacillus strain LB33

MELKRHYKMYKAGKKWVFAA IATISIIAGLNTVAVTYYAA
 GNNDPQQTTTQONAPNNSNDP QSTTTQNTANNSNDPQSTTT
 QNTANNSNGPQSTTTQNTAN NSNGPQSTTTQNTANNSNDP
 QSTTTQNTANNSNDPQSTTT QNTANNSNGPQSTTTQNTAN
 NSNDPQSTTTQNTANNGNDP QSTTGKDTVSIADIQVNPV 200
 NLLGKQSTVSSTGYNDSHIK NVNGKIYFVGDNQVKKNFT
 AIINGQSLYFNKTTGELASN DVQYENGLVKINDVHNAAYS
 IDP?GFTNVNGFLTANSWYR PKYIYKDGQKWVESTSQDMR
 PLLMTWVPDKNTQVAYLQYM QKMGILPADVTISSQTNQSV
 LTKESFITQAEIEKQIGVTN GNTDWLKKDISDFVNSQPNW 400
 NIDSEAKGTDHLQGGALLYV NNKLTPTYANS DYRLNRLTLT
 NQQGQVKDTSKQGGYEMLLA NDVDNSNPVQAEQLNWLYY
 MMNIGSITANDPTANFDGYR VDAVDNVDADLLNIAADYAK
 DAYKTNQSDANANKHLSILE DWDNNDPAYIKAHGNNQLTM
 DFPALHAIKYSINMPVSQRS GLEPELTTSLVNRTGDDSTE 600
 NVAQPNYTFIRAHDSVQTI IAQIIKDKINPNSDGLTVTP
 DEISQAFKIYNADLKTDKQ YTFYNMPSAYTILLTNKDTV
 PRVYYGDLYSDNGNYMSAHS PYYDAITLLKTRMKYVSGG
 QNMRMQYMQGDDMPANSYKG VLTŚVRYGKGEMTADEQGNS
 ETRTQIGIVIIISNNPNLKL DNDQVVLNMGAHENQTYRP 800
 VLLTTKDGKLYSDSDSSVPQ NALVSTNDKGQLIFKASSIQ
 GVSNPQVSGYLSVWVPVGA DNQDARTASSSQPSTDGKTY
 HSNAALDSQVIYEGFSNFQS IPTNTEDFTNVKIAQNANLF
 KSLGITSFELAPQYRSSNDN SFLDSVVQNGYAFTRDYDIG
 YNTPTKYGTVTQLLDALRAL HANGIQAIDDWVPDQIYNLP 1000
 GEEIVAAQRTNGSGTYDQDS VIDDTLYDSHTVGGGEYQAK
 FGGAFNLKQLYPDLFKVK QISTGQPMNPNERITEWSAK
 YFNGTNIQGRGAWYVLKDWG TNQYFNVSNNQFVPKQFLGT
 DTYTGFNVTNEGTFYSTSG YKAQNTFIQDGDNWYFDNN
 GYMVTGLQNINGNNYFLPN GIELQDSYLLNDDTGKEYYY 1200
 ASNGKQISNRYYPDANGNWR YFFNDGSMARNGLTITIEQPD

21/28

GQKVIQYFSDSGIQLKGNAA KDNNGNLRYPDGNTGDMVIN
SFGELPDGSWLYLNDKGI AV TGKQEINGQTYFADADGKQV
KNDFRELDPDGSWLYLNDKGI AVTG?QEINGQTYFADADGK
QVKNDRELDPDGSWLYLNDK GIAVTGKQGINGQTYAEAKI 1400
TAAENAHQAATDAVNKAQAA QSPNTSSSSSSSVSQATKHQL
AVKTAKAQLAKT?AQIAKYQ KALKKAKTTKAKAQARKSLK
KAETSFSKAELNLALFK

SEQ ID No. 19 DNA

SEQ ID No. 20 PRT

Lactobacillus sake strain KG15

1 SASCTGBCMSTNACGTTHRRCNTAGACGTTHRACGTACTGGTTCACACAATGGATTCGGC
1 X X X X R X X X T X X V L V H T M D S A

61 AAACATCAATGATTGCGATCTGTCCAGGTTGGGCTGCTTCACGCGTCAAACCGTACGG
21 N Y Q * L R S V Q V G L L H A S N Q Y G

121 ATCGCATTGACCACGGGTAATAATTGTAGTGC GCGACGGTTGAACCGTGACCGACTAATG
41 S H * P R V I I V V R D G * T V T D * W

181 GTGATTTTTTTCGGCATAAAGCGGTCATCAAGCGCCAAAAACGGCGTTGTGATTGAATA
61 * F F A A * R R S S S A K N G V V I E Y

241 CCAAGCGTTGTTTGTAAACACAGTAGCGCCAACAATCGACAGTCATCGATTTTAACGTGC
81 Q A L F V N T V A P T I D S H R F * R A

301 GCCACATTACGCCGTTGCGTCACACAACGTGGGCAATAGCGCTGGTAAAGCGACTGGCAC
101 P H Y A V A S H N V G N S A G K A T G T

361 AGCTGATAAAAAATAATGATAGTTACCTTGTAATTCGTGACGAATTTGTTTAACTTAGGA
121 A D K N N D S Y L V I R D E F V * T * D
- 35

421 TGGTTCAACATCGTTAGGACCCCTTTTAAAGTTTAGTCACTTATGAATCTAACTGTGTTGG
141 G S T S L G P L L S L V T Y E S N C V G
- 10

481 ACTTTTTTGTAAATTTTTTTTGTATTATTACAACTAGCACCACGCGTATGTGTTTATTA
161 L F C * F F C I I T N * H H A Y V F Y *
RBS

541 ATACCACCTTAATTAATAACGGGGCTTTAGCATGATTTCAAATAAAATAGTGTGAAAGGTA
181 Y H L I N N G A L A * F Q I K * C E R *
start

601 GTTTTTTATGTTAAGGAATAATTATTTTGGAGAGACTAAAACGCATTATAAATTATATAA
201 F F M L R N N Y F G E T K T H Y K L Y K

661 ATGCGGTAAGAACTGGGCTGTCATGGGGATTTTATTATTTCCGCTGGGATTAGGGATGCT
221 C G K N W A V M G I S L F P L G L G M L

721 AGTTACCAGCCAGCCAGTGTGCTGATGTGACAGCCACCAGCACCTCAAGCAGTGCAGT
241 V T S Q P V S A D V T A T S T S S S A V

781 GAGGACCGATGCAATCAGTGCAAGTAGTAGCAGTGCAGCAAAGGCTGAAACGGCTGCGAT
261 R T D A I S A S S S S A A K A E T A A I

841 CACTACTGCAGGTGTTGCAAATGCTGATTACAAACATCAGCAGAAGTAACCGCTGACTC
281 T T A G V A N A D S Q T S A E V T A D S

901 TACTTCTACCAGCCAAGTGGTAACTAATAATTCCAATAATCAAAATAATACAGCACAGCC
301 T S T S Q V V T N N S N N Q N N T A Q P

961 AGCCGGTCAAGAAGCAGCCCCGGTATCAGAGGACACATCATCTGATGATAGTGAGAGAAC
321 A G Q E A A P V S E D T S S D D S E R T

1021 TACACCAACAGTTGCAAATAATGATAAGCCAGCAATTGATTTCAGTTGACACTTCACAACC
341 T P T V A N N D K P A I D S V D T S Q P

1081 TGCAACTGCAGCGCCAAAAGCAGACACTGATGTATCAACGCTACAAGTAGATGCAACTAC
361 A T A A P K A D T D V S T L Q V D A T T

1141 GAAGACCGATTTCAGACATAAAAGAGGATACACCAACAGATAAGACAACCGATACAAAGAC
381 K T D S D I K E D T P T D K T T D T K T

1201 TGTGCAATTAACCACTGTTGAAGGAACGTCCAAGCAAGTGGTAACGACGCCGAAGGAAGA
401 V Q L T T V E G T S K Q V V T T P K E E

1261 GAGCTCAACTGACAAATCTTCGTCTGTGGTTTCTAAACAAACAGACAAAACGTCTTTGCC
421 S S T D K S S S V V S K Q T D K T S L P

1321 AACCGTAGCAACAGCAACAGCGACGACAGTGTCTAAGATTCCTTCAGTGACAGGTGATTA
441 T V A T A T A T T V S K I P S V T G D Y

1381 CCAGTTTGACGAAAAGACGAAGACTTATACGTTTACAGGTAAAGATGGTCATCCCCTAAC
461 Q F D E K T K T Y T F T G K D G H P V T

1441 TGGGTTGGTTTACGCGAATAATATCCTGCAATACTTTGATGAAACGGGTCATCAAGTAA
481 G L V Y A N N I L Q Y F D E T G H Q V K

1501 AGGTCAATACGTTACAATTGCAGGTCATGTATATTATTTTCGACCCAGCCAGCGCGCTGC
501 G Q Y V T I A G H V Y Y F D P A S G A A

1561 ACAAACAGGTGTTAATCAAATCGATGGTAAGATGGTTGGGTTTAAATCTGATGGGTCACA
521 Q T G V N Q I D G K M V G F K S D G S Q

1621 AATTACGTCAGGTTTTTCTAATGATAACGCCGAAATTCTTACTACTTTGATGAGTCTGG
541 I T S G F S N D N A G N S Y Y F D E S G

1681 AACCATGGTGACAGGGTGGCAAACCTATTGCTGGTAAGACGTATTACTTTGACAAAGACGG
561 T M V T G W Q T I A G K T Y Y F D K D G

1741 GCATCTCCGTAAGGGGTATTCCACTATTATTGATAATCAATTGTACTATTTTCGATTTGAA
581 H L R K G Y S T I I D N Q L Y Y F D L K

1801 AACAGGAGAGTCTGTTTCAACAACGACGTCCAATTTCAAATCTGGCTTGACATCACAAC
601 T G E S V S T T T S N F K S G L T S Q T

1861 GGATGACACAACACCACATAATAGTGCGGTTAATATGTCTAAGGATAGTTTTACCACCGT
621 D D T T P H N S A V N M S K D S F T T V

1921 TGATGGATTCTTGACAGCTGAGTCATGGTATGTACCTAAAGATATTCAAACATCTGCGAC
641 D G F L T A E S W Y V P K D I Q T S A T

1981 GGACTGGCGTGCATCAACGCCTGAAGATTTCCGTCCGATCATGATGACTTGGTGGCCAAC
661 D W R A S T P E D F R P I M M T W W P T

2041 GAAGCAAATTCAAGCAGCGTATTTGAACCATATGGTCTCCGAAGGATTGTTGTCATCAGA
681 K Q I Q A A Y L N H M V S E G L L S S D

2101 TAAGAAGTTCTCCGCAACGGATGATCAAACGTTGTTGAACCAAGCAGCACACGCGGTTCA
701 K K F S A T D D Q T L L N Q A A H A V Q

2161 ATTGCAAATTGAATTGAAGATTCAACAGACAAAGTCTGTTGAATGGTTGCGAACAACGAT
721 L Q I E L K I Q Q T K S V E W L R T T M

2221 GCACAAATTCATTAAAGTCACAACCAGGATACAATGTTACTAGTGAAACGCCAAGTAACGA
741 H N F I K S Q P G Y N V T S E T P S N D

2281 CCACCTTCAAGGTGGCGCATTAAAGCTACATTAACAGTGTGTTTGACGCCTGATGCGAACTC
761 H L Q G G A L S Y I N S V L T P D A N S

2341 AAATTTCCGTTTGATGAACCGTAATCCAACACAACAAGATGGTACGCGTCATTACAACAC
781 N F R L M N R N P T Q Q D G T R H Y N T

2401 TGATACATCTGAGGGTGGATATGAGTTGCTGTAGCCAATGACGTGGATAATTCTAACCC
801 D T S E G G Y E L L L A N D V D N S N P

2461 AGTTGTTCAAGCAGAACAATTGAACTGGTTGTACTTCTTGACGCATTTCCGGTGAAATTGT
821 V V Q A E Q L N W L Y F L T H F G E I V

2521 TAAGAACGATCCGTCAGCTAACTTTGATAGTGTTAGAGTGGATGCGGTAGACAACGTGGA
841 K N D P S A N F D S V R V D A V D N V D

2581 TGCCGACCTGCTAAACATTACAGCCGCTTATTTTAGAGATGTGTATGGCGTCGATAAAAA
861 A D L L N I T A A Y F R D V Y G V D K N

2641 CGATTTGACAGCTAATCAACATTTGTCTATTTTGAAGATTGGGGCCACAATGACCCATT
881 D L T A N Q H L S I L E D W G H N D P L

2701 ATATGTCAAGGACCACGGTAGTGATCAGTTGACGATGGATGATTACATGCATACCCAATT
901 Y V K D H G S D Q L T M D D Y M H T Q L

2761 GATTTGGTCATTAACAAAAAATCCAGATAATCGTAGTGCGATGCGACGATTTATGGAGTA
921 I W S L T K N P D N R S A M R R F M E Y

2821 TTATTTGGTCGACCGTGCTAAGGACAATACGTCTGATCCAGCAATTCCTAATTACAGCTT
941 Y L V D R A K D N T S D P A I P N Y S F

2881 TGTCCGTGCACACGATAGTGAAGTTCAAACGGTTATCGGTGATATTGTTGCGAAGTTGTA
961 V R A H D S E V Q T V I G D I V A K L Y

2941 TCCGGATGTTAAAAATTCATTGCCATCTATGGAACAATTGGCGGCAGCCTTTAAGGTATA
981 P D V K N S L P S M E Q L A A A F K V Y

3001 CGATGCGGATATGAATTCTGTTAATAAGAAGTATACGCAATACAACATGCCCGCAGCGTA
1001 D A D M N S V N K K Y T Q Y N M P A A Y

3061 TGCCATGTTACTAACGAATAAAGACACAATTCCACGTGTTTACTATGGTGATATGTATAC
1021 A M L L T N K D T I P R V Y Y G D M Y T

3121 GGATGATGGTCAATATATGGCAACTAAGTCACCATATTACGATGCCATCTCAGCGTTGTT
1041 D D G Q Y M A T K S P Y Y D A I S A L L

3181 GAAAGCCCGTATTAAGTATGTGGCTGGTGGACAAACCATGGCTGTAGATAAACACGATAT
1061 K A R I K Y V A G G Q T M A V D K H D I

3241 CTTAACATCAGTTCGCTTTGGTGATGGGATCATGAATGCATCTGATAAGGGTAGCACGAC
1081 L T S V R F G D G I M N A S D K G S T T

3301 GGCCCGTACCCAAGGAATTGGCGTGATTGTCAGCAATAATGATGCGTTAGCGTTGAAGGG
1101 A R T Q G I G V I V S N N D A L A L K G

3361 AGACACTGTGACCCCTTCATATGGGTATCGCTCACGCCAACCAGGCATACCGTGCTTTGTT
1121 D T V T L H M G I A H A N Q A Y R A L L

3421 GTTAACGACGACAGATGGACTAATGAAATACACGTCCGATAATGGCGCGCCAATTTCGTA
1141 L T T T D G L M K Y T S D N G A P I R Y

3481 TACGGATGCAAATGGTGACTTGATTTTCACTAGCGCAGACATTAAGGGATACAAAACGT
1161 T D A N G D L I F T S A D I K G Y Q N V

3541 TGAGGTATCCGATTCTTGTCTAGTGTGGGTACCAGTCGGTGCATCCGACACACAGGATGC
1181 E V S G F L S V W V P V G A S D T Q D A

3601 GCGCGCAACAGGGTCTAGCGCTGCAAACAAAACCTGGTGACACCTTACATTCAAATGCAGC
1201 R A T G S S A A N K T G D T L H S N A A

3661 ATTGGACTCAAATGTGATTTATGAAGGTTTTCTAATTTCCAAGAGATGCCAACAGCCCA
1221 L D S N V I Y E G F S N F Q E M P T A H

3721 CGATGAGTTTACAAACGTAAAGATCGCTCAAAATGCTGATTTGTTTAAAGTCATGGGGTGT
1241 D E F T N V K I A Q N A D L F K S W G V

3781 GACAAGTTTCCAACCTTGACCCACAATATCGTTCAAGTGATGACACATCATTTTTGGATTCT
1261 T S F Q L A P Q Y R S S D D T S F L D S

3841 TATTATTAAGAATGGATATGCGTTTACAGACCGCTATGACTTGGGCTTTAATACGCCAAC
1281 I I K N G Y A F T D R Y D L G F N T P T

3901 AAAGTACGGAGACGTTGACGACCTAGCAGATGCAATTAGAGCAATGCACAGTGTGGTAT
1301 K Y G D V D D L A D A I R A M H S V G I

3961 TCAGGTCATGGCTGACTTTGTCCCTGACCAAATTTATAATTTGCCAGGTCAAGAAGTAGT
1321 Q V M A D F V P D Q I Y N L P G Q E V V

4021 TGCTGTTAATCGTACTAATAACTTTGGTACACCAAACCAGGATTCAGATCTACAAAACCA
1341 A V N R T N N F G T P N Q D S D L Q N Q

4081 GTTGATGTTACAAATTCAAAGGGTGGCGGTGAATACCAAGCTAAGTATGGTGGTGAGTT
1361 L Y V T N S K G G G E Y Q A K Y G G E F

4141 CTTGGATCTTTTTCGCTCTGGAACACCCTGATTTGTTTACAACAAATCAGATTTTCGACTGG
1381 L D L L R L E H P D L F T T N Q I S T G

4201 TGTACCAATCGATGGGTCCACGAAGATTAAAGAATGGTCTGCAAAGTACTTCAATGGTTC
1401 V P I D G S T K I K E W S A K Y F N G S

4261 TGACATCCAAGGTAAGGGCGCTGATTACGTATTGAAGGATGGTGCATCTCAAGAATATTT
1421 D I Q G K G A D Y V L K D G A S Q E Y F

4321 CAAGATTACGTCTAATGCGAACGATGAGTCCTTCTTGCCAAAACAATTTATGAATCAAGA
1441 K I T S N A N D E S F L P K Q F M N Q D

4381 TGCCATGACTGGGTTCACCACAGATGAAAAGGGCACAACCTTATTATTCAACAAGTGGTTA
1461 A M T G F T T D E K G T T Y Y S T S G Y

4441 CCAAGCTAAACAGTCGTTTATCCAAGGTGATGATGGACAATATTATTACTTTGATGCAGA
1481 Q A K Q S F I Q G D D G Q Y Y Y F D A D

4501 CGGATACATGGTGCAGGGCTCACAAACCATTAATGGTAAGCAATATTACTTCTTGCCAAA
1501 G Y M V T G S Q T I N G K Q Y Y F L P N

4561 TGGCGTTGAGTTAAGAGAAGCATTTTTACAAAATGCATCTGGTAACACGGTTTATTATGG
1521 G V E L R E A F L Q N A S G N T V Y Y G

4621 CAAGACTGGCTCAGCAGTTAAGTCTAAATATGTAGTCGATCAAAGCGGTGTGGCTTATTA
1541 K T G S A V K S K Y V V D Q S G V A Y Y

4681 CTTTGATGTAAACGGTAATATGGTTCGAGATCGTATGATGATTCTTGATGGACACACGCA
1561 F D V N G N M V A D R M M I L D G H T Q

4741 ATATTTCTTTGCGGGTGGTTTCAAGCTAAGGACCAATTTTTGATTGGGTCAGATGGTAA
1581 Y F F A G G S Q A K D Q F L I G S D G N

4801 CTTACGTTACTTTGACCAAGGTAGTGGTAATATGGTTACAAATCGTTTTGCAGTAAACCG
1601 L R Y F D Q G S G N M V T N R F A V N R

4861 AAACGGGGATTGGTTCTACTTCAATGGCGATGGTATCGCGTTGAAGGGTTGGCAAACAAT
1621 N G D W F Y F N G D G I A L K G W Q T I

4921 TGCTGGTAAGACTTATTTCTTTGATGCTGATGGACGTCAAGTCAAGGCTGCCGCTGACAA
1641 A G K T Y F F D A D G R Q V K A A A D K

4981 GGCTGCTGCTGAACAAGCCGCTGCTGACAAGGCTGCCGCTGAACAAGCCGCTGCTGACAA
1661 A A A E Q A A A D K A A A E Q A A A D K

5041 GGCTGCCGCTAAGGATAAGCAAACCTCAAGCTGTAGCTTACGCTGCTACCAAGGCTAAGAA
1681 A A A K D K Q T Q A V A Y A A T K A K N

5101 CAATATTGATCAAGCTACTACAGCTGATGGCATCAATGATGCCCAAGCAACTGGTATCAC
1701 N I D Q A T T A D G I N D A Q A T G I T

5161 TGATATTGATAACCAGCATGTTCTGTTGATAATAAAAAGCAAGCTGAGAA
1721 D I D N Q H V P G T S V D N K K Q A E K

5221 GGTAAGTGAAGATATCAAGAATGATCCAGATAATAAGACTTTGCCTGAAGCTATCGAATT
1741 V T E D I K N D P D N K T L P E A I E L

5281 ACCAAATACGGGCGTTGATAAGACAGAAAGTATTACTATTACCGGTGTAGTTATGCTAAT
1761 P N T G V D K T E S I T I T G V V M L I
stop

5341 CCTCACTACTATTTTTGGTCTGTTGTTTACAAGTAAAAAGCATAAAAAGATTAGTGTAG
1781 L T T I F G L L F T S K K H K K D * C R

5401 ATAGCTATACCAAAGGGAGTTAACATAACATCGATTATTGAGATATGAAGTTATTTAGGG
1801 * L Y Q R E L T * H R L F R Y E L I * G
-----> inverted

5461 ACTATAATTTACAAATAACCCCTATGCAACGCTATTTAAACAACCCCGTTATCTATTGG
1821 L * F T N N P Y A T L L K Q P P L S I G
repeat <----- (-10.7 Kcal mol⁻¹)

5521 ACAGGTAATAGGGTTGTTTTATGTTTTTTATGGCAGATTGCAAGAAATAACTTGAAC
1841 Q V I G V V F M F F Y G R L Q E I T * T

5581 AAATTTAGTAACGCAGATTACGCAAAAAGATCTCAATCGGTGTTGCGCAATTTAAACATT
1861 N L V T Q I T Q K D L N R C S P I * T F

5641 TGAGTGGTCGGGAATTCAAATACCAGTTAATTTGAATCAATTCATCATCGGTAATCTCAT
1881 E W S G I Q I P V N L N Q F I I G N L I

5701 CAATCGGTTGACCTTTGGGAATGAAGCGGCGTAGTACTCGGTTGCGATTTTCATTACTGC
1901 N R L T F G N E A A * Y S V A I F I T A

5761 CTCGTTTCATGTGGCGAATAAGCGTGCGCAAAGTACAGCGGAACACCGGCTGTTCTTCAA
1921 S F M W R I S V R K V Q R N T G L F F N

5821 TCAAATTGTAGTTGGCAAATTCTTTCCCATGGTCAACAGTAAGGGTCTTGAGATTATCTC
1941 Q I V V G K F F P M V N S K G L E I I S

5881 CTAAGTGGTTAGCCAATTCTAAAATAGCCTTGGTCATTGAAGTACTATCTCGTCCATTAA
1961 * L V S Q F * N S L G H * S T I S S I K

5941 GCCGTTTAACGATGGTAAGCCGACTCTTACGCTCGACAAACGTAGCTACTGCTTGACCTT
1981 P F N D G K P T L T L D K R S Y C L T F

6001 TACGTTTTCCAGATAAAACTGTATCAGCTTCGAAGTGACCTGAAGTATTACGATCAGAAA
2001 T F S R * N C I S F E V T * S I T I R N

6061 TTTCAGCTGGACGATCTTCGATGGAACGTCCATGACTAAAACTACCACGGGTTTCTTTAG
2021 F S W T I F D G T S M T K T T T G F F S

6121 CTCGTTTTCGACGAATACCATGGTCAGGTAAATCAGTCACATTAAT
2041 S F A T N T M V R * I S H I N

SEQ ID No. 21 DNA

SEQ ID No. 22 PRT

Lactobacillus fermentum strain LB33

1 ATTAATGGCCGCATTTGTTGTGACACAGCCACAGTGGAATAAAACAAGTGAAGATGTGAA
1 L M A A F V V T Q P Q W N K T S E D V N

61 TGATGATCATTTGCAAGGTGGGGCATTAAACATTTGAAAAATAACGGCGACACAGACGCTAA
21 D D H L Q G G A L T F E N N G D T D A N

121 TTCGGATTATCGCCTCATGAACCGCACGCCAACAAATCAGACTGGCGAACGCTTGTACCA
41 S D Y R L M N R T P T N Q T G E R L Y H

181 CATTGATGACTCACTTGGTGGTTACGAATTATTGCTGGCAAATGACGTTGACAATTCAAA
61 I D D S L G G Y E L L L A N D V D N S N

241 TCCACAAGTTCAGGCAGAACAATTGAATTGGTTGTACTACTTAATGCATTTTGGGGATAT
81 P Q V Q A E Q L N W L Y Y L M H F G D I

301 TACAGCTGATGATCCGGACGCAAATTTTGATGCCATACGGATTGATGCGGTGCGATAATGT
101 T A D D P D A N F D A I R I D A V D N V

361 CGATGCTGATTTACTTCAACTAGCAGCCCAGTATTTCCGGGATGCCTATGGCATGGCTAC
121 D A D L L Q L A A Q Y F R D A Y G M A T

421 AACTGACGCAACATCAAATAAGCATCTTCAATTCTTGAGGATTGGAGCCATAACGATCC
141 T D A T S N K H L S I L E D W S H N D P

481 GCGGTATATGCAAGCACACGGCAATGATCAATTAACGATGGATGATTATATGCACACACA
161 A Y M Q A H G N D Q L T M D D Y M H T Q

541 GTTGATTTGGTCATTAACCAAGCCCAGGCACAACGCGGGACCATGGCACGCTTTATGGA
181 L I W S L T K P E A Q R G T M A R F M D

601 CTTCTATCTCACCAACCGTGCTAATGATGATACAGAAAACACGGCGCAACCTAGTTACTC
201 F Y L T N R A N D D T E N T A Q P S Y S

661 GTTTGTGCGTGCCCATGATAGCGAAGTACAAACAGTCATTGCTGAGATCGTGACGAAGCT
221 F V R A H D S E V Q T V I A E I V T K L

721 GCATCCAGAAGCAGGAAATGGGTAAATGCCTACGGAAGAACAAATGGCAGAAGCGTTTAA
241 H P E A G N G L M P T E E Q M A E A F K

781 GATTTACAATGCGGACCAAAAGAAGGCCGTTAAGACTTACACACATTACAATATGCCATC
261 I Y N A D Q K K A V K T Y T H Y N M P S

841 TGCATACGCCATGCTGTAAACGAACAAGGATGTTATTCCACGAATTTACTATGGTGACTT
281 A Y A M L L T N K D V I P R I Y Y G D L

901 GTACACTGATGATGGGCAATTCATGGCGACAAAATCACCTTATTTTGATGCGATTTTCGAC
301 Y T D D G Q F M A T K S P Y F D A I S T

961 CATGTTACAAGCACGCACGAAGTATGTAGCTGGTGGACAGACGATGGCGGTTGACCAGCA
321 M L Q A R T K Y V A G G Q T M A V D Q H

1021 CGACGTCTTGACTAGCGTTCGGTTTTGGTAAGGGGGCCATGACGGCCAATGATTTAGGGGA
341 D V L T S V R F G K G A M T A N D L G D

1081 TGCTGAGACCCGGACTGAGGGTGTGGGATTAATTATTAGCAACAACCCAAAGTTGCAATT
361 A E T R T E G V G L I I S N N P K L Q L

1141 GGGACAACAAGACAACGTGGTGTACACATGGGACTTGCGCACGCGAATCAGGCATTCCG
381 G Q Q D N V V L H M G L A H A N Q A F R

1201 CGCAGTCGTACTAACGACCGCGACCGGATTAACCATTTATAATGACGATGATGCTCCGAT
401 A V V L T T A T G L T I Y N D D D A P I

1261 TCGTTATACCGATAATAAGGGTGATTTAATTTTCACTAACCATGACGTATATGGCGTGT
421 R Y T D N K G D L I F T N H D V Y G V L

1321 GAATCCACAAGTGTGTCAGGCTTCTTGCCAATGTGGGTGCCAACTGGTGCACCAGCGAACCA
441 N P Q V S G F L A M W V P T G A P A N Q

1381 GGATGCGCGATCTACTGCGTCAACCAACATGTCAACGGATGGATCTGCCTACCATTCTAA
461 D A R S T A S T N M S T D G S A Y H S N

1441 TGCGGCTTTGGATAGTCAAGTAATCTTTGAATCATTTCGAATTTCCAGGCTATGCCAAC
481 A A L D S Q V I F E S F S N F Q A M P T

1501 AAGTCATGACACATACACCAACGTTGTGTAGCCAATCATGCTGACCAGTTGCACGATTG
501 S H D T Y T N V V L A N H A D Q L H D W

1561 GGGAATAACTTCGGTACAGTTAGCACCACAATACCGGTCTTCAACCGACGGTACCTTTTT
521 G I T S V Q L A P Q Y R S S T D G T F L

1621 AGACGCGATTATTCAAATGGCTATGCCTTCACTGACCGTTATGATTTAGGGTTTGGTAC
541 D A I I Q N G Y A F T D R Y D L G F G T

1681 GCCAACTAAATACGGGGATGATACGGATTTGCGGAACGTCATCAAAGCATTGCATGCAAA
561 P T K Y G D D T D L R N V I K A L H A N

1741 TGGCATGCAAGTAATGGCTGATTTTGTGCCGGATCAATTGTATACATTACCAGGTAAGGA
581 G M Q V M A D F V P D Q L Y T L P G K E

1801 ATTGGTACAAGTCACCCGAACAAACAATATGGGTGAGCCAGATACGCATTCTGACATCCA
601 L V Q V T R T N N M G E P D T H S D I Q

1861 ACATATTTTATATGTGACGAGCACTCGTGGTGGTGGTGACTATCAGAAACAGTACGGTGG
621 H I L Y V T S T R G G G D Y Q K Q Y G G

1921 TGAGTTCCTTGACGATTGCGTGAACGATACCCAGATTTATTTACGACACGTCAAATTTTC
641 E F L A R L R E R Y P D L F T T R Q I S

1981 GACCGGACAAACAATTGATGATTGAGTAAATAAAGAATGGTCAGCTAAGTATTTGAA
661 T G Q T I D D S V K I K E W S A K Y L N

2041 TGGTACCGCAATTCAAGGACGTGGAGCTGGCTATGTGCTGCGTGATAATGGTACAAATGC
681 G T A I Q G R G A G Y V L R D N G T N A

2101 TTATTACAAGGTGACAGCAAATGACGGTAATGTGAACTTACCAAAGCAATTACTCGGCCA
701 Y Y K V T A N D G N V N L P K Q L L G Q

2161 ACCGGTGATGACCGGATTCTATCACGAGGCAGATGGTTATCATTTTGAAACATTGAGTGG
721 P V M T G F Y H E A D G Y H F E T L S G

2221 TACGTCGGCCAAAGATGCCTTTATTATGGGCGACGATGGGGCACTGTATTATTTTGATGA
741 T S A K D A F I M G D D G A L Y Y F D D

2281 TCAGGGTGTATTATGGTAACGGGTAAGCAACGTGTGCACCAAGATCAGTATTTCTTCCTGCC
761 Q G V M V T G K Q R V H Q D Q Y F F L P

2341 AAATGGTATTGCTTTGACAGATGCTTTTCGTACAAACTGCTGATGGTCAACGTCAGTACTA
781 N G I A L T D A F V Q T A D G Q R Q Y Y

2401 TGATAAAACAGGTCGTCTGGTCATTAATCAATATGTGACTGACCACCAAGCGAATGCGTT
801 D K T G R L V I N Q Y V T D H Q A N A F

2461 CCGGGTTGATGCAGACGGTAACGTTGTCCGCAATCAAGCTTTGACTGTTGACGGCCATGA
821 R V D A D G N V V R N Q A L T V D G H E

2521 ACAATATTTTCGGCACAAACGGTGTCCAAGCGAAAGCAGTGCTCATTTCGAACTGACGATAA
841 Q Y F G T N G V Q A K A V L I R T D D N

2581 TCAGGCGCGCTACTACGAAGCCAATAGTGGTAATCTCGTGAAGCAACAGTTTATTCTTGA
861 Q A R Y Y E A N S G N L V K Q Q F I L D

2641 TACAGATGGACATTGGTTGTACGCGGATGCTGCAGGTGACTTGGCACGCGGACAAATTAC
881 T D G H W L Y A D A A G D L A R G Q I T

2701 AATTGGCCAAGACACGTTGTATTTTGATGATAATAATCACCAGGTAAAAGATGATTTTCGT
901 I G Q D T L Y F D D N N H Q V K D D F V

2761 CTATGATACTAACGGTGTGCATTATTTTAATGGCACAAACAGGCGCTGAAATCAAACAAGA
921 Y D T N G V H Y F N G T T G A E I K Q D

2821 TTACGCGTTTCATGATGGCAAATGGTACTATTTTGATGATTTGGGACGAATGGTAACCGG
941 Y A F H D G K W Y Y F D D L G R M V T G

2881 CTTGCAGCGTATTAATGGTGAGTATCGCTATTTTGATGCTAATGGTGTGCAACTAAAGGG
961 L Q R I N G E Y R Y F D A N G V Q L K G

2941 CCGGTACCGTGACCGATCCACTAACGCACCAACGTACACTTTTGATGCGAAAACCTGGTGC
981 G T V T D P L T H Q T Y T F D A K T G A

3001 TGGTACGTTGGTGACGATTTAACTGAATAATGGACTAGAAAAGACGATCTTGTATCGTCT
1001 G T L V T I * L N N G L E K T I L Y R L

3061 TTTTAgTTTCGATAACTAAATAAGTGCTCATTTTTCATTAGGACTCAGAATTAGCGGG
1021 F * F R * L N K C S F L H * D S E L A G

3121 CGCGCAAGCGTCTTTTCGTGTTAACTTATTAGTAATTAATATTTTGAGGAGTCTGTTAT
1041 A Q A S F R V K L I S N * Y F E E S V I

3181 ATGGCAACAATTTTAGTTGTAGATGATGAACCGTCATTGGTGACGCTACTGTCATAACAAC
1061 W Q Q F * L * M M N R H W * R Y C H T T

3241 CTGACTAAATCAGGCTTCGAGGTCGTGACTGCTACCTCCGGTGACGAGGCACGAAATCAG
1081 * L N Q A S R S * L L P P V T R H E I S

3301 CTGGCAAATCATCCTATTGATTTGATGCTGCTAGGTGTCATGTTGCCTGGTAAGAGTGGC
1101 W Q I I L L I * C C * V S C C L V R V A

3361 GTTGACTTAACACGAGAACTACGAGGCGAACAGAATCGTATTCCAATTATTATGATTACC
1121 L T * H E N Y E A N R I V F Q L L * L P

3421 GCCTTGGATGACGAAGTTGACAAGATTT
1141 P W M T K L T R F

(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization
International Bureau



(43) International Publication Date
30 January 2003 (30.01.2003)

PCT

(10) International Publication Number
WO 03/008618 A3

(51) International Patent Classification⁷: **C12P 19/18**,
19/08, C12N 15/52, 9/10, 1/20, 1/21, C08B 37/02, A23L
1/054 // (C12P 19/18, C12R 1:225)

(74) Agent: **JORRITSMA, Ruurd**; Nederlandsch Octrooi-
bureau, Scheveningseweg 82, P.O. Box 29720, NL-2502 LS
The Hague (NL).

(21) International Application Number: **PCT/NL02/00495**

(81) Designated States (*national*): AE, AG, AL, AM, AT, AU,
AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU,
CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,
GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC,
LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW,
MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG,
SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ,
VN, YU, ZA, ZM, ZW.

(22) International Filing Date: **22 July 2002 (22.07.2002)**

(25) Filing Language: **English**

(26) Publication Language: **English**

(30) Priority Data:
01202752.0 20 July 2001 (20.07.2001) EP
01202841.1 25 July 2001 (25.07.2001) EP

(84) Designated States (*regional*): ARIPO patent (GH, GM,
KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW),
Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM),
European patent (AT, BE, BG, CH, CY, CZ, DE, DK, EE,
ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK,
TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GQ,
GW, ML, MR, NE, SN, TD, TG).

(71) Applicant (*for all designated States except US*): **NED-
ERLANDSE ORGANISATIE VOOR TOEGEPAST-
NATUUR-WETENSCHAPPELIJK ONDERZOEK
TNO [NL/NL]**; Schoemakerstraat 97, NL-2628 VK Delft
(NL).

Published:
— *with international search report*

(72) Inventor; and

(88) Date of publication of the international search report:
30 October 2003

(75) Inventor/Applicant (*for US only*): **VAN GEEL-SCHUT-
TEN, Gerritdina, Hendrika [NL/NL]**; Bosstraat 78,
NL-3971 XG Driebergen (NL).

*For two-letter codes and other abbreviations, refer to the "Guid-
ance Notes on Codes and Abbreviations" appearing at the begin-
ning of each regular issue of the PCT Gazette.*

(54) Title: **GLUCANS AND GLUCANSUCRASES DERIVED FROM LACTIC ACID BACTERIA**

(57) Abstract: The invention pertains to glucans capable of being produced by glucosyltransferase activity of a lactic acid bacterium on a sucrose substrate, the glucan having an average molecular weight between 10 kDa and 1 GDa, consisting essentially of alpha (1,3)- and alpha (1,6)-linked anhydroglucose units (AGU) and to glucansucrases capable of producing these glucans from sucrose. The glucans have thickening and anti-corrosive properties. The glucans can be chemically modified.

WO 03/008618 A3

INTERNATIONAL SEARCH REPORT

International Application No
PCT/NL 02/00495

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 C12P19/18 C12P19/08 C12N15/52 C12N9/10 C12N1/20
C12N1/21 C08B37/02 A23L1/054 //(C12P19/18,C12R1:225)

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 C12P C08B C12N

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, WPI Data, BIOSIS, SEQUENCE SEARCH, PAJ

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	VAN GEEL-SCHUTTEN G H ET AL: "Screening and characterization of Lactobacillus strains producing large amounts of exopolysaccharides." APPLIED MICROBIOLOGY AND BIOTECHNOLOGY, vol. 50, no. 6, December 1998 (1998-12), pages 697-703, XP002233876 ISSN: 0175-7598 the whole document --- -/--	1-6, 10-14, 16-20

☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

* Special categories of cited documents:

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier document but published on or after the international filing date
- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

- "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- "&" document member of the same patent family

Date of the actual completion of the international search

7 March 2003

Date of mailing of the international search report

18. 06. 2003

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl
Fax: (+31-70) 340-3016

Authorized officer

Madruga, J

INTERNATIONAL SEARCH REPORT

Internation No
PCT/NL 02/00495

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	<p>UZOCHUKWU SYLVIA ET AL: "Structural analysis by ¹³C-nuclear magnetic resonance spectroscopy of glucans elaborated by gum-producing bacteria isolated from palm wine."</p> <p>FOOD CHEMISTRY, vol. 73, no. 2, May 2001 (2001-05), pages 225-233, XP002233877 ISSN: 0308-8146 page 233, left-hand column, paragraph 1 - paragraph 2; figure 3; tables 1,2</p> <p>---</p>	1-5,14, 16-20
X	<p>PIDOUX M ET AL: "MICROSCOPIC AND CHEMICAL STUDIES OF A GELLING POLYSACCHARIDE FROM LACTOBACILLUS-HILGARDII" CARBOHYDRATE POLYMERS, vol. 13, no. 4, 1990, pages 351-362, XP002233878 ISSN: 0144-8617 the whole document</p> <p>---</p>	1-4,14, 16-20
X	<p>MONCHOIS V ET AL: "Cloning and sequencing of a gene coding for a novel dextransucrase from Leuconostoc mesenteroides NRRL B-1299 synthesizing only alpha(1-6) and alpha(1-3) linkages" GENE: AN INTERNATIONAL JOURNAL ON GENES AND GENOMES, ELSEVIER SCIENCE PUBLISHERS, BARKING, GB, vol. 182, no. 1-2, 5 December 1996 (1996-12-05), pages 23-32, XP004071926 ISSN: 0378-1119 page 27, paragraph 3 page 29, paragraph 2 Conclusions page 31</p> <p>---</p>	3,4, 10-13, 16-20
X	<p>ARGUEELLO-MORALES M A ET AL: "SEQUENCE ANALYSIS OF THE GENE ENCODING ALTERNANSUCRASE, A SUCROSE GLUCOSYLTRANSFERASE FROM LEUCONOSTOC MESENEROIDES NRRL B-1355" FEMS MICROBIOLOGY LETTERS, AMSTERDAM, NL, vol. 182, 2000, pages 81-85, XP000937860 ISSN: 0378-1097 the whole document</p> <p>---</p>	3,4, 10-13, 16-20
X	<p>US 5 789 209 A (COTE GREGORY L ET AL) 4 August 1998 (1998-08-04) column 1, line 21 - line 52; figures column 3, line 18 - line 29 column 4, line 2 - line 111; examples</p> <p>---</p>	3,4,10, 16-20

-/--

INTERNATIONAL SEARCH REPORT

Intern: Application No
PCT/NL 02/00495

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	VUYST DE L ET AL: "HETEROPOLYSACCHARIDES FROM LACTIC ACID BACTERIA" FEMS MICROBIOLOGY REVIEWS, ELSEVIER, AMSTERDAM, NL, vol. 23, no. 2, 1999, pages 153-177, XP000971896 ISSN: 0168-6445 page 169, left-hand column, paragraph 2 -page 170, left-hand column, paragraph 1 ---	1-6, 10-14, 16-20
A	PATENT ABSTRACTS OF JAPAN vol. 018, no. 468 (C-1244), 31 August 1994 (1994-08-31) & JP 06 146036 A (NIPPON SYNTHETIC CHEM IND CO LTD:THE;OTHERS: 01), 27 May 1994 (1994-05-27) abstract ---	19
A	PATENT ABSTRACTS OF JAPAN vol. 2000, no. 02, 29 February 2000 (2000-02-29) & JP 11 310895 A (SUMITOMO METAL IND LTD), 9 November 1999 (1999-11-09) abstract ---	19
A	EP 0 427 349 A (TNO) 15 May 1991 (1991-05-15) cited in the application the whole document -----	16

INTERNATIONAL SEARCH REPORT

International application No.
PCT/NL 02/00495

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:

2. ☐ Claims Nos.:
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:

3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

see additional sheet

1. ☐ As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.

2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.

3. ☐ As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:

4. ☒ No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

6 (completely); 1-5, 10-14, 16-20 (all partially)

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.

This International Searching Authority found multiple (groups of) inventions in this international application, as follows:

1. Claims: 6 completely. Claims 1-5, 10-14 and 16-20, all in part

A glucan capable of being produced by a glucosyltransferase activity of a lactic acid bacteria on a sucrose substrate, the glucan having an average molecular weight of 50 KDa-1 MDa, and having a backbone consisting of alpha(1,3)- and alpha(1,6)-linked anhydroglucose units (AGU), comprising 30-45% of alpha(1,3)-linked AGU, 30-45% of alpha(1,6)-linked AGU and 3-13% of alpha (1,3, 6)-linked AGU; a chemically modified glucan; uses of said glucan.

A Lactobacillus strain producing said glucan (Lb33).

A glucosyltransferase enzyme from Lactobacillus able to produce said glucan, nucleic acid encoding therefore (corresponding to Lb33, SEQ ID NOs: 3,4,17,18), host cells containing said nucleic acid and process to produce said glucan.

2. Claims: 7 completely. Claims 1-5, 10-14 and 16-20, all in part

A glucan capable of being produced by a glucosyltransferase activity of a lactic acid bacteria on a sucrose substrate, the glucan having an average molecular weight of 10-50 MDa, and having a backbone consisting of alpha(1,3)- and alpha(1,6)-linked anhydroglucose units (AGU), comprising 12-26% of alpha(1,3)-linked AGU, 30-50% of alpha(1,6)-linked AGU and 5-20% of alpha (1,3, 6)-linked AGU; a chemically modified glucan; uses of said glucan.

A Lactobacillus strain producing said glucan (strain 180).

A glucosyltransferase enzyme from Lactobacillus able to produce said glucan, nucleic acid encoding therefore (corresponding to strain 180, SEQ ID NOs: 1,2, 11, 12), host cells containing said nucleic acid and process to produce said glucan.

3. Claims: 1-5, 8, 10-14 and 16-20, all in part

A glucan capable of being produced by a glucosyltransferase activity of a lactic acid bacteria on a sucrose substrate, the glucan having an average molecular weight of 1-50 MDa, and having a backbone consisting of alpha(1,3)- and

alpha(1,6)-linked anhydroglucose units (AGU) (comprising 45-60% of alpha(1,3)-linked AGU, 4-10% of alpha(1,6)-linked AGU and 10-20% of alpha (1,3,6)-linked AGU; a chemically modified glucan; uses of said glucan.

A Lactobacillus strain producing said glucan (L. reuterii ML1).

A glucosyltransferase enzyme from Lactobacillus able to produce said glucan, nucleic acid encoding therefore (corresponding to L. reuterii ML1, SEQ ID NOs: 13,14), host cells containing said nucleic acid and process to produce said glucan.

4. Claims: 1-5, 8, 10-14 and 16-20, all in part

A glucan capable of being produced by a glucosyltransferase activity of a lactic acid bacteria on a sucrose substrate, the glucan having an average molecular weight of 1-50 MDa, and having a backbone consisting of alpha(1,3)- and alpha(1,6)-linked anhydroglucose units (AGU) (comprising 45-60% of alpha(1,3)-linked AGU, 4-10% of alpha(1,6)-linked AGU and 10-20% of alpha (1,3,6)-linked AGU; a chemically modified glucan; uses of said glucan.

A Lactobacillus strain producing said glucan (L. reuterii ML4).

A glucosyltransferase enzyme from Lactobacillus able to produce said glucan, nucleic acid encoding therefore (corresponding to L. reuterii ML4, SEQ ID NOs: 15,16), host cells containing said nucleic acid and process to produce said glucan.

5. Claims: 1-4, 9, 10-14 and 16-20, all in part

A glucan capable of being produced by a glucosyltransferase activity of a lactic acid bacteria on a sucrose substrate, the glucan having an average molecular weight of 10-50 MDa, and having a backbone consisting of alpha(1,3)- and alpha(1,6)-linked anhydroglucose units (AGU) (comprising 80-99% of alpha(1,6)-linked AGU and 0-15% of alpha(1,3)-linked AGU; a chemically modified glucan; uses of said glucan.

A Lactobacillus strain producing said glucan (strain LB 33).

A glucosyltransferase enzyme from Lactobacillus able to produce said glucan, nucleic acid encoding therefore (corresponding to strain LB 33, SEQ ID NOs: 17,18), host cells containing said nucleic acid and process to produce said glucan.

6. Claims: 1-4, 9, 10-14 and 16-20, all in part

A glucan capable of being produced by a glucosyltransferase activity of a lactic acid bacteria on a sucrose substrate, the glucan having an average molecular weight of 10-50 MDa, and having a backbone consisting of alpha(1,3)- and alpha(1,6)-linked anhydroglucose units (AGU) (comprising 80-99% of alpha(1,6)-linked AGU and 0-15% of alpha(1,3)-linked AGU; a chemically modified glucan; uses of said glucan.

A Lactobacillus strain producing said glucan (L. sake KG15).

A glucosyltransferase enzyme from Lactobacillus able to produce said glucan, nucleic acid encoding therefore (corresponding to L. sake KG15, SEQ ID NOs: 19,20), host cells containing said nucleic acid and process to produce said glucan.

7. Claims: Claims 9-20 (all in part)

A glucosyltransferase enzyme from Leuconostoc able to produce a glucan on a sucrose substrate, a glucan having an average molecular weight of 10-50 MDa, and comprising 88-99% of alpha(1,6)-linked AGU, a nucleic acid encoding therefor (corresponding to Lc 86-1, partial sequence SEQ ID NO: 3), host cells containing said nucleic acid and process to produce said glucan.

A Leuconostoc strain producing said glucan.

8. Claims: Claims 9-20 (all in part)

A glucosyltransferase enzyme from Leuconostoc able to produce a glucan on a sucrose substrate, a nucleic acid encoding therefor (corresponding to Lc 86-5, partial sequence SEQ ID NO: 7, SEQ ID NO: 8), host cells containing said nucleic acid and process to produce said glucan.

9. Claims: Claims 9-20 (all in part)

A glucosyltransferase enzyme from Leuconostoc able to produce a glucan on a sucrose substrate, a nucleic acid encoding therefor (corresponding to Lc 86-8, partial sequence SEQ ID NO: 9, SEQ ID NO: 10), host cells containing said nucleic acid and process to produce glucan.

INTERNATIONAL SEARCH REPORT

Inte Application No

PCT/NL 02/00495

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
US 5789209	A	04-08-1998	AU 697819 B2	15-10-1998
			AU 3235295 A	04-03-1996
			CA 2196596 A1	15-02-1996
			EP 0804545 A1	05-11-1997
			WO 9604365 A1	15-02-1996
			US 5702942 A	30-12-1997

JP 06146036	A	27-05-1994	NONE	

JP 11310895	A	09-11-1999	NONE	

EP 0427349	A	15-05-1991	NL 8902786 A	03-06-1991
			NL 9001027 A	18-11-1991
			AT 124960 T	15-07-1995
			CA 2029542 A1	11-05-1991
			DE 69020861 D1	17-08-1995
			DE 69020861 T2	30-11-1995
			DK 427349 T3	20-11-1995
			EP 0427349 A2	15-05-1991
			ES 2074528 T3	16-09-1995
			GR 3017196 T3	30-11-1995
			JP 3172171 B2	04-06-2001
			JP 4175301 A	23-06-1992
			AT 174346 T	15-12-1998
			AU 649038 B2	12-05-1994
			AU 7696091 A	27-11-1991
			CA 2081483 A1	28-10-1991
			DE 69130608 D1	21-01-1999
			DE 69130608 T2	27-05-1999
			DK 526494 T3	16-08-1999
			EP 0526494 A1	10-02-1993
			ES 2126570 T3	01-04-1999
			FI 924843 A	26-10-1992
			GR 3029515 T3	28-05-1999
			JP 3004052 B2	31-01-2000
			JP 5506685 T	30-09-1993
			KR 204437 B1	15-06-1999
			WO 9117189 A1	14-11-1991
			NO 924134 A	27-10-1992
			US 5326864 A	05-07-1994
